



US009482671B2

(12) **United States Patent**  
**Gennaro**(10) **Patent No.:** **US 9,482,671 B2**  
(45) **Date of Patent:** **\*Nov. 1, 2016**

- (54) **PROTEINS EXPRESSED BY MYCOBACTERIUM TUBERCULOSIS AND NOT BY BCG AND THEIR USE AS DIAGNOSTIC REAGENTS AND VACCINES**
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- (\*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 0 days.
- This patent is subject to a terminal disclaimer.

(21) Appl. No.: **14/989,135**(22) Filed: **Jan. 6, 2016**(65) **Prior Publication Data**

US 2016/0153986 A1 Jun. 2, 2016

**Related U.S. Application Data**

- (60) Division of application No. 14/201,308, filed on Mar. 7, 2014, now Pat. No. 9,238,066, which is a division of application No. 13/893,659, filed on May 14, 2013, now Pat. No. 8,974,800, which is a division of application No. 13/198,108, filed on Aug. 4, 2011, now Pat. No. 8,992,942, which is a continuation of application No. 12/503,717, filed on Jul. 15, 2009, now Pat. No. 8,021,832, which is a continuation of application No. 11/677,502, filed on Feb. 21, 2007, now Pat. No. 7,579,141, which is a division of application No. 10/009,383, filed as application No. PCT/US00/12257 on May 4, 2000, now Pat. No. 7,932,373.
- (60) Provisional application No. 60/132,505, filed on May 4, 1999.

(51) **Int. Cl.**

**A61K 39/04** (2006.01)  
**A61K 39/02** (2006.01)  
**A61K 39/00** (2006.01)  
**G01N 33/569** (2006.01)  
**C07K 14/35** (2006.01)  
**G01N 33/50** (2006.01)  
**A61K 38/00** (2006.01)

(52) **U.S. Cl.**

CPC ..... **G01N 33/5695** (2013.01); **A61K 39/04** (2013.01); **C07K 14/35** (2013.01); **G01N 33/5091** (2013.01); **A61K 38/00** (2013.01); **A61K 39/00** (2013.01); **A61K 2039/53** (2013.01); **G01N 2333/35** (2013.01); **G01N 2333/57** (2013.01); **G01N 2800/26** (2013.01); **Y10S 435/863** (2013.01)

(58) **Field of Classification Search**

CPC ..... **A61K 38/00**; **A61K 39/00**; **A61K 39/04**; **G01N 33/5695**  
 USPC ..... 424/184.1, 185.1, 234.1, 248.1; 435/7.1, 7.2, 253.1

See application file for complete search history.

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**ABSTRACT**

The present invention is directed to reagents useful for generating immune responses to *Mycobacterium tuberculosis* and for diagnosing infection and disease in a subject that has been exposed to *M. tuberculosis*.

**8 Claims, 8 Drawing Sheets**

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MTBN1

MTAEPEVRTLREVVLDDQLGTAESRAYKMWLPPLTNPVPLNELIARDRRQPLRFALGIMDE  
PRRHLODVWGVVDVSGAGGNIGIGGAPQTGKSTLLQTMVMSAAATHSPRNVOFYCIDLGGG  
GLIYLENLPHVGGVANRSEPDKNRVVAEMQAVMRQRETTFKEHVRVGSIGMYRQLRDDPS  
QPVASDPYGDVFLIIDGWFGFVGEFFDLEGQVQDLAAQGLAFGVHVIISTPRWTELKSRV  
RDYLGTKIEPRLGDVNETQIDRITREIPANRPFRAVSMEXHMLIGVPRFDGVHSADNLV  
EAITAGVTQIASQHQTEQAPPVRVLPERIHLEHDPNPPGPESDYRTRWEIPIGLRETDLT  
PAHCHMHTNPHLLIFGAAGSGKTTIAHAIAARAICARNSPQQVRFMLADYRSGLLDVDPDT  
HLLGAGAINRNSASLDEAVQALAVNLKKRLPPTDLTTAQLRSRSWWSGPDVVLLVDDWHM  
IVGAAGGMPMPAPLAPLLPAAADIGLHIIVTCQMSQAYKATMDKPFVGAAPFGSGAPTMFLS  
GEKQEFPSSEFKVKRRPFGQAFVSPDGKEVIQAPYIEPPERVFAPPSAG

MTBN2

MEKMSHDPAAADIGTQVSDNALHGVTAGSTALTSVTGLVPAGADEVSAQAATAFTSEGIO  
LLASNASAQDQLHRAGEAVQDVARTYSOIDDGAAGVFAE

MTBN3

MLWHAMPPELNTARLMAGAGPAPMLAAAAGWQTLAALDAQAVELTARLNSLGEAWTGGG  
SDKALAAATPMVVWLQTAQTAKTRAMQATAQAAAYTQAMATTPSLPEIAANHITQAVLT  
ATNFFGINTIPIALTEMDYFIRMWNQAALAMEVYQAEAVNTLFEKLEPMASILDPGASQ  
STTNPIFGMPSPGSSSTFVQQLPFAATOTLGLGEMSGPMQQLTQPLQQVTSLSFSQVGGTG  
GQNPADDEEAAQMGLLGTSPLSNHPLAGGSGPSAGAGLLRAESLPGAGGSLTRTPILMSQLI  
EKPVAPSVMPAAAAGSSATGGAAPVGAGAMGQGAQSGGSTRPGLVAPAPLAQEREDEDD  
DWDEEDDW

MTBN4

MAEMKTDAATLAQEAGNFERISGDLKTQIDQVESTAGSLQQQWRGAAGTAAQAAVVRFQE  
AANKQKQELDEISTNIRQAGVQYSRADEEQQALSSQMGF

MTBN5

MAADYDKLFRPHEGMEAPDDMAAQPFDPASAFPPAPASANLPKPNGQTFPPPTSDDLSE  
FVSAPPPPPPPPPPPPTPMPIAAGEPPSTEPAASKPPTPPMPIAGPEPAPPKPPTPEMP  
IAGPEPAPPKPPTPPMPIAGPAPTPTESQLAPPRPPTPQTPTGAPQQPESPAPHVPSHGP  
HQPRRTAPAPPWAKMPIGEPPEAPSRPSASPAEFPTRPAPQHSRRARRGHRVRTDTERNV  
GKVATGPSIQARLRAEEASGAQLAPGTEPSAPFLGQPRSYLAPFTRPAPTEPPPSPSQOR  
NSGRRRAERRVHPDLAAQHAAAQPSITAATTGGRARRKRAAPDLDTQKSLRPAKGPVK  
KVKPQKPKATKPPKVVSQRGWRHWHALTRINLGLSPDEKYELDLHARVRNRPRGSYQIA  
VVGLKGGAGKTTTLTAALGSTLAQVRADRIALADADPGAGNLADRVGRQSGATTADVLAEK  
ELSHYNDIRAHTSVNAVNLVLPAPEYSSAQRALSDADWHFIADPASRFYNLVLADCGAG  
FFDPLTRGVLSTVSGVVVVASVSDGAQQASVALDWLRNNGYQDLASRACVVINHIMPGE  
PNVAVKDLVRHFEQQVQPGRVVMPWDRHIAAGTEISLDLLDPYKRVLELAAALSDDF  
ERAGRR

FIG 1A

MTBN6

LSAPAVAAGPTAAGATAARPATTRVTILTGRRM TD LVLPAAVPMETYIDDTVAVLSEVLE  
DTPADVLGGFDFTAQGVWAFARFGSPPLKLDQSLDDAGVVDGSLLTLSVSRTERYRPLV  
EDVIDAIAVLDESPEFDRTALNRFVGAAPLLTAPVIGMAMRAWWETGRSLWWPLAIGIL  
GIAVLVGSFVANRFYQSGHLAECLLVTTYLLIATAAALAVPLPRGVNSLGAPQVAGAATA  
VLFLLTLMTRGGPRKRHELASFVITAIAVIAAAAAFGYGYQDWVPAGGIAFGLFIVTNA  
KLTVAVARIALPPIPVPGETVDNEELLDPVATPEATSEETPTWQAIIASVPASAVRLTER  
SKLAKQLLIGYVTSGLTILAAGAI VVVVRGHFFVHSLVVAGLITTVCGFRSRLYAERWCA  
WALLAATVAIPTGLTAKLIWYPHYAWLLLSVYLTVALVALVVVGSMAHVRRVSPVVKRT  
LELIDGAMIAAIPMLLWITGVYD TVRNIRF

MTBN7

MAEPLAVDPTGLSAAAAKLAGLVFPQPPAPIAVSGTDSVVAAINETMPSIESLSVDGLPG  
VKAALTRTASNMNAAADVAKTDQSLGTSLSQYAFGSSGEGLAGVASVGGQPSQATQLLS  
TPVSQVTTQLGETAAELAPRVVATVPQLVQLAPHAVQMSQNASPIAQTISQTAQQAQSA  
QGGSGPMPAQLASAEKPATEQAEPVHEVTNDDQGDQGDVQPAEVVAAARDEGAGASPGQQ  
PGGGVPAQAMDTGAGARPAASELAAPVDPSTPAPSTTTTL

MTBN8

MSITRPTGSYARQMLDPGGWVEADEDTFYDRAQEYSQVLQRVTDVLDTCRQQKGHVFEGG  
LWSGGAANAANGALGANINQLMTLQDYLATVITWHRHIAGLIEQAKSDIGNNV DGAQREI  
DILENDPSLDADERHTAINSLVTATHGANVSLVAETAERVLESKNWKPPKNALEDLLQOK  
SPPPPDVPTLVVPSFGTPGTPTGTPTITPGTPTITPGTPTITPIPGAPVTPITPTPGTPVTPVT  
PGKPVTPVTPVKPGTPGEPTPTITPTVTPFVAPATPATPATPVTPAPAPHPQAPAPAPSPG  
PQPVTPTATPGPSGPATPGTPGGEPAHPVKPAALAEQPGVPGQHAGGQTQSGPAHADESAA  
SVTPAAASGVPGARAAAAAPSGTAVGAGARSSVGTAAASGAGSHAATGRAPVATSDKAAA  
PSTRAASARTAPPARPPSTDHIDKPDRESADDGTPVSMIPVSAARAARDAATAAASARQ  
RGRGDALRLARRIAAALNASDNNAGDYGFFWITAVTTDGSIVVANSYGLAYIPDGMELPN  
KVYLASADHAIPVDEIARCATYPVLAVQAWAAFHDMTLRAVIGTAEQLASSDPGVAKIVL  
EPDDIPESGKMTGRSRLEVVDPSAAAQLADTTDQRLDLLPPAPVDVNP PGDERHMLWFE  
LMKPMTSTATGREAAHLRAFRAYAHSQEIALHQAHTATDAAVQRVAVADWLYWQYVTGL  
LDRALAAAC

FIG 1B

mtbn1

1 atgactgctg aaccggaagt acggacgctg cgcgagggtg tgotggacca  
51 gctcggcact gctgaatcgc gtgcgtacaa gatgtggctg ccgccgttga  
101 ccaatccggt cccgctcaac gagctcatcg cccgtgatcg gcgacaaccc  
151 ctgcgatttg ccttggggat catggatgaa ccgcgccgcc atctacagga  
201 tgtgtggggc gtagacgttt ccggggcccg cggcaacatc ggtattgggg  
251 gcgcacctca aaccgggaag tcgacgctac tgcagacgat ggtgatgtcg  
301 gccgccgcca cacactcacc gcgcaacggt cagttctatt gcatcgacct  
351 aggtggcggc gggctgatct atctcgaaaa ccttcacacg gtccgtgggg  
401 tagccaatcg gtccgagccc gacaagggtc accgggtggg cgcagagatg  
451 caagccgtca tgcggcaacg ggaaaccacc ttcaaggaac accgagtggg  
501 ctcgatcggg atgtaccggc agctgcgtga cgatccaagt caaccggtg  
551 cgtccgatcc atacggcgac gtctttctga tcatcgacgg atggcccggt  
601 tttgtcggcg agttccccga ccttgagggg caggttcaag atctggccgc  
651 ccaggggctg gcgttcggcg tccacgtcat catctccacg ccacgctgga  
701 cagagctgaa gtgcgctggt cgcgactacc tcggcaccaa gatcgagttc  
751 cggcttggtg acgtcaatga aaccagatc gaccggatta cccgcgagat  
801 cccggcgaat cgtccgggtc gggcagtgct gatggaaaag caccatctga  
851 tgatcggcgt gccaggttc gacggcgtgc acagcgccga taacctggtg  
901 gaggcgatca ccgcgggggt gacgcagatc gcttcccagc acaccgaaca  
951 ggcacctcgc gtgcgggtcc tgcggagcgc tatccacctg caogaactcg  
1001 accgaaccc gccgggacca gactccgact accgcactcg ctgggagatt  
1051 ccgatcggct tgcgcgagac ggacctgacg ccggctcact gccacatgca  
1101 caggaacccg cacctactga tcttcggtgc ggccaaatcg ggcaagacga  
1151 ccattgccca cgcgatcgcg cgcgccattt gtgcccgaaa cagtccccag  
1201 caggtgcggg tcatgtctgc ggactaccgc tcgggcctgc tggacgcggt  
1251 gccggacacc catctgctgg gcgccggcgc gatcaaccgc aacagcgctg  
1301 cgctagacga ggccgttcaa gcaactggcg tcaacctgaa gaagcggttg  
1351 ccgccgaccg acctgacgac ggcgcagcta cgtcgcggt cgtggtggag  
1401 cggatttgac gtcgtgcttc tggtcgacga ttggcacatg atcgtgggtg  
1451 ccgccggggg gatgccgcgc atggcacgcg tggccccgtt attgccggcg  
1501 gcggcagata tcgggttgca catcattgtc acctgtcaga tgagccaggc  
1551 ttacaaggca accatggaca agttcgtcgg ccgcccatte gggtcgggcg  
1601 ctccgacaat gttcctttcg ggcgagaagc aggaattccc atccagtga  
1651 ttcaagggtc agcggcgccc ccttgccag gcattttctc tctcgccaga  
1701 cggcaaagag gtcattccag cccctacat cgagcctcca gaagaagtgt  
1751 tcgcagcacc cccaagcgcc ggttaa

mtbn2

1 atggaaaaaa tgtcacatga tccgatcgct gccgacattg gcacgcaagt  
51 gagcgacaac gctctgcacg gcgtgacggc cggtcgcacg gcgctgacgt  
101 cggtgaccgg gctgggtccc gcgggggccc atgaggtctc cgcccaagcg  
151 gcgacggcgt tcacatcgga gggcatccaa ttgctggctt ccaatgcac  
201 ggcccaagac cagctccacc gtgcgggcga agcggtcag gacgtcgccc  
251 gcacctattc gcaaatcgac gacggcgccc ccggcgtctt cgccgaatag

FIG 2A

mtbn3

1 atgctgtggc acgcaatgcc accggagcta aataccgcac ggctgatggc  
51 cggcgcggt cgggtccaa tgcttgccgc ggccgcggga tggcagacgc  
101 tttcggcggt tctggacgct caggccgtcg agttgaccgc gcgcctgaac  
151 tctctgggag aagcctggac tggagggtggc agcgacaagg cgcttgccgc  
201 tgcaacgccg atggtggtct ggctacaaac cgcgtcaaca caggccaaga  
251 cccgtgcgat gcaggcgacg gcgcaagccg cggcatacac ccaggccatg  
301 gccacgacgc cgtcgtgcc ggagatcgcc gccaaaccaca tccccaggc  
351 cgtccttacg gccaccaact tcttcggtat caacacgac ccgatcgct  
401 tgaccgagat ggattatttc atccgtatgt ggaaccaggc agccctggca  
451 atggaggtct accaggccga gaccgcggtt aacacgcttt tcgagaagct  
501 cgagccgatg gcgtcgatcc ttgatcccg cgcgagccag agcacgacga  
551 acccgatctt cggaatgccc tccctggca gctcaacacc ggttggccag  
601 ttgccgcggc cggtaccca gacctcggc caactgggtg agatgagcgg  
651 cccgatgcag cagctgacct agccgctgca gcaggtgacg tcgttgttca  
701 gccaggtggg cggcaccggc ggcggcaacc cagccgacga ggaagccgcg  
751 cagatgggccc tgctcggcac cagtccgctg tcgaaccatc cgctggctgg  
801 tggatcaggc cccagcgccg gcgcgggcct gctgcgcgcg gactcgctac  
851 ctggcgaggg tgggtcggtg acccgcacgc cgtgatgtc tcagctgac  
901 gaaaagccgg ttgccccctc ggtgatgccg gcggctgctg ccgatcgctc  
951 ggcgacgggt ggcgcgcgtc cggtggtgctc gggagcgatg ggccagggtg  
1001 cgcaatccgg cggctccacc aggcggggtc tggtcgcgcc ggcaccgctc  
1051 gcgcaggagc gtgaagaaga cgacgaggac gactgggacg aagaggacga  
1101 ctggtga

mtbn4

1 atggcagaga tgaagaccga tgccgctacc ctgcgcagg aggcaggtaa  
51 tttcgagcgg atctccggcg acctgaaaac ccagatcgac cagggtggagt  
101 cgacggcagg ttcgttgacg ggccagtgcc gcggcgccgc ggggacggcc  
151 gccaggccg cggtggtgct cttccaagaa gcagccaata agcagaagca  
201 ggaactcgac gagatctcga cgaatatctg tcaggccggc gtccaatact  
251 cgagggccga cgaggagcag cagcaggcgc tgtcctcgca aatgggcttc  
301 tga

mtbn5

1 atggcgggccg actacgacaa gctcttcggc ccgcacgaag gtatggaagc  
51 tccggacgat atggcagcgc agccgttctt cgaccccagt gcttcgtttc  
101 cgccggcgcc cgcacggca aacctaccga agcccaacgg ccagactccg  
151 ccccgacgt ccgacgacct gtcggagcgg ttcgtgctcg ccccgccgc  
201 gccaccccca ccccaacctc cgctccgccc aactccgatg ccgatcgccg  
251 caggagagcc gccctcgccg gaaccggccg catctaaacc acccacacc  
301 cccatgcca tcgccggacc cgaaccggcc ccacccaaac caccacacc  
351 ccccatgccc atcgccggac ccgaaccggc cccacccaaa ccaccacac  
401 ctccgatgcc catcgccgga cctgcaccca ccccaaccga atcccagttg

FIG 2B

451 gcgcccccca gaccaccgac accacaaacg ccaaccggag cgccgcagca  
501 accggaatca ccggcgcccc acgtaccctc gcacgggcca catcaacccc  
551 ggcgcacccg accagcaccg ccctgggcaa agatgccaat cggcgaaccc  
601 ccgcccgcgc cgtccagacc gtctgcgtcc ccggccgaac caccgacccg  
651 gcctgcccc caacaotccc gacgtgcgcg ccggggtcac cgctatcgca  
701 cagacaccga acgaaacgtc ggggaaggtag caactggtcc atccatccag  
751 gcgcggctgc gggcagagga agcatccggc gcgcagctcg ccccggaac  
801 ggagccctcg ccagcgccgt tgggccaaacc gagatcgtat ctggctccgc  
851 ccacccgccc cgcgccgaca gaacctcccc ccagcccctc gccgcagcgc  
901 aactccggtc ggctgcccga gcgacgcgtc caccocgatt tagccgccc  
951 acatgccgcg gcgcaacctg attcaattac ggccgcaacc actggcggtc  
1001 gtcgccgcaa gcgtgcagcg ccgatctcgc acgcgacaca gaaatcctta  
1051 aggcggcgcg ccaagggggc gaagggtgaag aagggtgaagc ccagaaaacc  
1101 gaaggccacg aagccgcccc aagtgggtgc gcagcgcggc tggcgacatt  
1151 ggggtcatgc gttgacgcga atcaacctgg gcctgtcacc cgacgagaag  
1201 tacgagctgg acctgcacgc tcgagtcgcg cgcaatcccc gcgggtcgta  
1251 tcagatcgcc gtcgtcggtc tcaaagggtg ggctggcaaa accacgctga  
1301 cagcagcggt ggggtcgacg ttggctcagg tgcgggcca cggatcctg  
1351 gctctagacg cggatccagg cgccggaaac ctgcgcgatc gggtagggcg  
1401 acaatcgggc gcgaccatcg ctgatgtgct tgcagaaaaa gaactgtgca  
1451 actacaacga catccgcgca cacactagcg tcaatgcggc caactggga  
1501 gtgctgccgg caccggaata cagctcggcg cagcgcgcg ctagcgacgc  
1551 cgactggcat ttcctgcgcg atcctgcgtc gaggttttac aacctcgtct  
1601 tggctgattg tggggccggc ttcttcgacc cgtgacccg cggcgtgctg  
1651 tccacgggtg ccggtgtcgt ggtcgtggca agtgtctcaa tcgacggcgc  
1701 acaacaggcg tcggtcgcgt tggactggtt gcgcaacaac ggttaccaag  
1751 atttggcgag ccgcgcacgc gtgggtcatc atcacatcat gccgggagaa  
1801 cccaatgtcg cagttaaaga cctgggtgcg catttcgaac agcaagttca  
1851 acccggcggc gtcgtggtca tgccgtggga caggcacatt gccggccgga  
1901 ccgagatttc actcgacttg ctcgacccta tctacaagcg caaggtcctc  
1951 gaattggccg cagcgctatc cgacgatttc gagagggctg gacgtcgttg  
2001 a

mtbn6

1 ttgagcgcac ctgctgttgc tgctggctct accgcccggc gggcaaccgc  
51 tgcgcggcct gccaccaccc ggggtgacgat cctgaccggc agacggatga  
101 ccgatttggg actgccagcg gcggtgccga tggaaactta tattgacgac  
151 accgtcgcgg tgctttccga ggtgttgga gacacgccg ctgatgtact  
201 cggcggttcc gactttaccg cgcaaggcgt gtgggcgttc gctcgtcccg  
251 gatcgccgcc gctgaagctc gaccagtcac tcgatgacgc cggggtggtc  
301 gacgggtcac tgctgactct ggtgtcagtc agtcgcaccg agcgtaccg  
351 accgttggtc gaggatgtca tcgacgcgat cgccgtgctt gacgagtcac  
401 ctgagttcga ccgcacggca ttgaatcgct ttgtgggggc ggcgatcccg  
451 cttttgaccg cgcccgtcat cgggatggcg atgcgggcgt ggtgggaaac  
501 tgggcgtagc ttgtgggtggc cgttggcgat tggcatcctg gggatcgtcg

FIG 2C



551 tgctggtagg cagcttcgtc gcgaacaggt tctaccagag cggccacctg  
601 gccgagtgcc tactgggtcac gacgtatctg ctgatcgcaa ccgccgcagc  
651 gctggccgtg ccgttgccgc gcgggggtcaa ctcgttgggg gcgccacaag  
701 ttgccggcgc cgttacggcc gtgctgtttt tgacctgat gacgcggggc  
751 ggccctcgga agcgtcatga gttggcgctg tttgccgtga tcaccgctat  
801 cgcgggtcatc gcggccggccg ctgccttcgg ctatggatac caggactggg  
851 tccccgcggg ggggatcgca ttccgggctgt tcattgtgac gaatgcggcc  
901 aagctgaccg tcgcgggtcgc gcggatcgcg ctgccgccga ttccgggtacc  
951 cggcgaaacc gtggacaacg aggagttgct cgatcccgtc gcgaccccg  
1001 aggtaccag cgaagaaacc ccgacctggc aggccatcat ccgctcgggtg  
1051 ccgcggtccg cgggtccggct caccgagcgc agcaaactgg ccaagcaact  
1101 totgatcgga tacgtcacgt cgggcaccct gattctggct gccggtgcc  
1151 tcgcggtcgt ggtgcgcggg cacttctttg tacacagcct ggtggtcgcg  
1201 ggtttgatca cgacggtctg cggatttcgc tcgcggcttt acgcgagcg  
1251 ctggtgtgcg tgggcgttgc tggcggcgac ggtcgcgatt ccgacgggtc  
1301 tgacggccaa actcatcatc tggtagccgc actatgctg gctgttgttg  
1351 agcgtctacc tcacggtagc cctggttgcg ctcggtgtgg tcgggtcgat  
1401 ggctcacgtc cggcgcgttt caccggctcg aaaacgaact ctggaattga  
1451 tcgacggcgc catgatcgct gccatcatte ccatgctgct gtggatcacc  
1501 ggggtgtacg acacgggtccg caatatccgg ttctga

mtbn7

1 atggtgaac cgttgccgt cgatcccacc ggcttgagcg cagcgccgcg  
51 gaaattggcc ggcctcgttt ttccgcagcc tccggcgccg atcgcggtca  
101 gcggaacgga ttccggtggtg gcagcaatca acgagaccat gccaaagcatc  
151 gaatcgctgg tcagtgaagg gctgcccggc gtgaaagccg cctgactcg  
201 aacagcatcc aacatgaacg cggcggcgga cgtctatgcg aagaccgatc  
251 agtcactggg aaccagtttg agccagtatg cattcggtc gtcgggcgaa  
301 ggccctggctg gcgtcgccctc ggtcggtggg cagccaagtc aggctacca  
351 gctgctgagc acacccgtgt cacagggtcac gacccagctc ggcgagacgg  
401 ccgtgagct ggcacccgt gttgttgoga cgggtgcgca actcgttcag  
451 ctggtccgc acgcggttca gatgtcgcaa aacgcacccc ccatcgctca  
501 gacgatcagt caaacgcgcc aacaggcgcg ccagagcgcg cagggcgga  
551 gcggcccaat gccgcacag cttgccagcg ctgaaaaacc ggccaccgag  
601 caagcggagc cgggtccacga agtgacaaac gacgatcagg gcgaccaggg  
651 cgacgtgcag ccggccgagg tcgttgccgc ggcacgtgac gaaggcgccg  
701 gcgcacacc gggccagcag ccgcggggg gcgttcccgc gcaagccatg  
751 gataccggag ccggtgcccg ccagcgggg agtccgctgg cggccccgt  
801 cgatccgtcg actccggcac cctcaacaac cacaacgttg tag

mtbn8

```
1 atgagtatta ccaggccgac gggcagctat gccagacaga tgctggatcc
51 gggcggctgg gtggaagccg atgaagacac tttctatgac cgggccagg
101 aatatagcca ggttttgcaa agggtcaccg atgtattgga cacctgccgc
151 cagcagaaag gccacgtctt cgaaggcggc ctatgggtccg gcggcgccgc
201 caatgctgcc aacggcgccc tgggtgcaaa catcaatcaa ttgatgacgc
251 tgcaggatta tctcgccacg gtgattacct ggcacaggca tattgccggg
301 ttgattgagc aagctaaatc cgatatcggc aataatgtgg atggcgctca
351 acgggagatc gatatcctgg agaatgaccc tagcctggat gctgatgagc
401 gccataccgc catcaattca ttggtcacgg cgacgcattg ggccaatgtc
451 agtctggctg ccgagaccgc tgagcgggtg ctggaatcca agaattggaa
501 acctccgaag aacgcactcg aggatttgc tcagcagaag tcgccgccac
551 ccccagacgt gcttacctg gtctgacct ccccgggcac accgggcaca
601 ccgggaaccc cgatcacccc gggaacccc atcaccccgg gaacccaat
651 cacaccatc ccgggagcgc cggtaactcc gatcacacca acgcccggca
701 ctcccgtcac gccggtgacc ccgggcaagc cggtcacccc ggtgaccccg
751 gtcaaaccgg gcacaccagg cgagccaacc ccgatcacgc cggtcacccc
801 ccgggtcgcc ccggccacac cggcaacccc ggccacgccc gttaccccag
851 ctcccgtccc acaccgcag ccggctccgg caccggcgcc atcgctggg
901 cccagccgg ttacaccggc cactcccgtt ccgtctggtc cagcaacacc
951 gggcacccca gggggcgagc cggcgccgca cgtcaaacc gcggcggttg
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1201 gcgggggtgc atgctgccac tgggcggggc ccggtggcta cctcggaaca
1251 ggcggcgga ccgagcacgc gggcggcctc ggcgcggacg gcacctctg
1301 ccgccccgc gtcgaccgat cacatcgaca aaccgatcg cagcgagtct
1351 gcagatgacg gtacgccggt gtcgatgate ccggtgtcgg cggctcgggc
1401 ggcacgcgac gccgccactg cagctgccag cgcgcgcag cgtggccgcg
1451 gtgatgcgct gcggttggcg cgacgcacgc cggcggcgct caacgcgtcc
1501 gacaacaacg cgggcgacta cgggttcttc tggatcaccc cggtgaccac
1551 cgacggttcc atcgtcgtgg ccaacagcta tgggctggcc tacatacccg
1601 acgggatgga attgccaat aaggtgtact tggccagcgc ggatcacgca
1651 atcccggttg acgaaattgc acgctgtgcc acctacccgg ttttggcgt
1701 gcaagcctgg gcgctttcc acgacatgac gctgcgggcg gtgatcggta
1751 ccgcgagca gttggccagt tcggatcccg gtgtggccaa gatttgtctg
1801 gagccagatg acattccgga gagcggcaaa atgacgggcc ggtcgcggt
1851 ggaggtcgtc gacccctcgg cggcggtcga gctggccgac actaccgata
1901 agcgtttgct cgacttgttg ccgcggcgcc cggtggtatg caatccaccg
1951 ggcgatgagc ggcacatgct gtggttcgag ctgatgaagc ccatgaccag
2001 caccgctacc ggccgcgagg ccgctcatct gcgggcgttc cgggcctacg
2051 ctgcccactc acaggagatt gcctgcacc aagcgcacac tgcgactgac
2101 cggcgctccc agcgtgtggc cgctcgcgac tggctgtact ggcaatacgt
2151 caccgggttg ctcgaccggg ccctggccgc cgcattgctga
```

FIG 2E

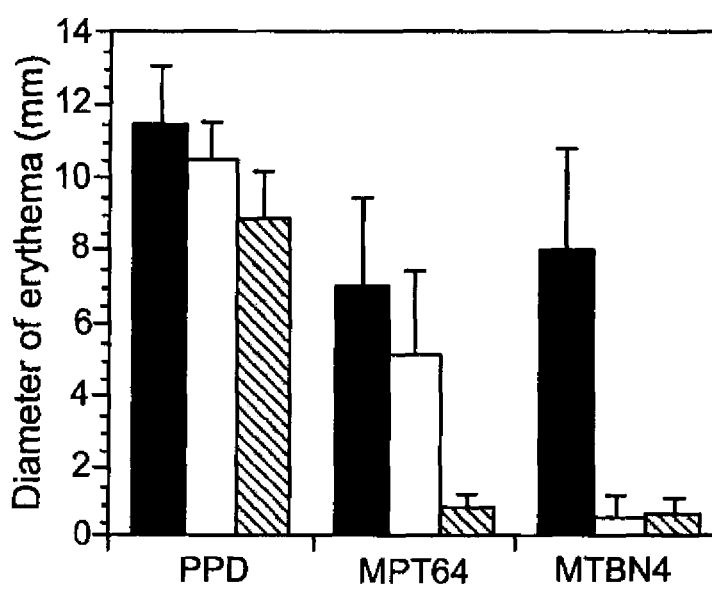


FIG 3

1

# **PROTEINS EXPRESSED BY MYCOBACTERIUM TUBERCULOSIS AND NOT BY BCG AND THEIR USE AS DIAGNOSTIC REAGENTS AND VACCINES**

This application is a Divisional of U.S. patent application Ser. No. 14/201,308 filed Mar. 7, 2014, now U.S. Pat. No. 9,238,066, which is a divisional of, and claims priority to, U.S. application Ser. No. 13/893,659, filed May 14, 2013, now U.S. Pat. No. 8,974,800, which is a divisional of, and claims priority to, U.S. application Ser. No. 13/198,108, filed Aug. 4, 2011, now U.S. Pat. No. 8,992,942, which is a continuation of, and claims priority to, U.S. application Ser. No. 12/503,717, filed Jul. 15, 2009 and now U.S. Pat. No. 8,021,832, which is a continuation of, and claims priority to, U.S. application Ser. No. 11/677,502, filed Feb. 21, 2007, now U.S. Pat. No. 7,579,141, which is a divisional of, and claims priority to, U.S. application Ser. No. 10/009,383, filed Mar. 4, 2002 and now U.S. Pat. No. 7,932,373, which claims priority to International Patent Application No. PCT/US00/12257, filed May 4, 2000, which claims priority to U.S. Provisional Application Ser. No. 60/132,505, filed May 4, 1999, the disclosures of each of which are hereby incorporated by reference in their entireties.

## **BACKGROUND OF THE INVENTION**

Tuberculosis infection continues to be a world-wide health problem. This situation has recently been greatly exacerbated by the emergence of multi-drug resistant strains of *M. tuberculosis* and the international AIDS epidemic. It has thus become increasingly important that effective vaccines against and reliable diagnostic reagents for *M. tuberculosis* be produced.

The disclosure of U.S. Pat. No. 6,087,163 is incorporated herein by reference in its entirety.

## **SUMMARY OF THE INVENTION**

The invention is based on the inventor's discovery that a polypeptide encoded by an open reading frame (ORF) in the genome of *M. tuberculosis* that is absent from the genome of the Bacille Calmette Guérin (BCG) strain of *M. bovis* elicited a delayed-type hypersensitivity response in animals infected with *M. tuberculosis* but not in animals sensitized with BCG. Thus proteins encoded by ORFs present in the genome of *M. tuberculosis* but absent from the genome of BCG represent reagents that are useful in discriminating between *M. tuberculosis* and BCG and, in particular, for diagnostic methods (e. g., skin tests and in vitro assays for *M. tuberculosis*-specific antibodies and lymphocyte responsiveness) which discriminate between exposure of a subject to *M. tuberculosis* and vaccination with BCG. The invention features these polypeptides, functional segments thereof, DNA molecules encoding either the polypeptides or the functional segments, vectors containing the DNA molecules, cells transformed by the vectors, compositions containing one or more of any of the above polypeptides, functional segments, or DNA molecules, and a variety of diagnostic, therapeutic, and prophylactic (vaccine) methodologies utilizing the foregoing.

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Specifically, the invention features an isolated DNA molecule containing a DNA sequence encoding a polypeptide with a first amino acid sequence that can be the amino acid sequence of the polypeptide MTBN1, MTBN2, MTBN3, MTBN4, MTBN5, MTBN6, MTBN7 or MTBN8, as depicted in FIGS. 1A and 1B, or a second amino acid sequence identical to the first amino acid sequence with conservative substitutions; the polypeptide has *Mycobacterium tuberculosis* specific antigenic and immunogenic properties. Also included in the invention is an isolated portion of the above DNA molecule. The portion of the DNA molecule encodes a segment of the polypeptide shorter than the full-length polypeptide, and the segment has *Mycobacterium tuberculosis* specific antigenic and immunogenic properties. Other embodiments of the invention are vectors containing the above DNA molecules and transcriptional and translational regulatory sequences operationally linked to the DNA sequence; the regulatory sequences allow for expression of the polypeptide or functional segment encoded by the DNA sequence in a cell. The invention encompasses cells (e.g. eukaryotic and prokaryotic cells) transformed with the above vectors.

The invention encompasses compositions containing any of the above vectors and a pharmaceutically acceptable diluent or filler. Other compositions (to be used, for example, as DNA vaccines) can contain at least two (e. g., three, four, five, six, seven, eight, nine, ten, twelve, fifteen, or twenty) DNA sequences, each encoding a polypeptide of the *Mycobacterium tuberculosis* complex or a functional segment thereof, with the DNA sequences being operationally linked to transcriptional and translational regulatory sequences which allow for expression of each of the polypeptides in a cell of a vertebrate. In such compositions, at least one (e. g., two, three, four, five, six, seven, or eight) of the DNA sequences is one of the above DNA molecules of the invention. The encoded polypeptides will preferably be those not encoded by the genome of cells of the BCG strain of *M. bovis*.

The invention also features an isolated polypeptide with a first amino acid sequence that can be the sequence of the polypeptide MTBN1, MTBN2, MTBN3, MTBN4, MTBN5, MTBN6, MTBN7 or MTBN8 as depicted in FIGS. 1A and 1B, or a second amino acid sequence identical to the first amino acid sequence with conservative substitutions. The polypeptide has *Mycobacterium tuberculosis* specific antigenic and immunogenic properties. Also included in the invention is an isolated segment of this polypeptide, the segment being shorter than the full-length polypeptide and having *Mycobacterium tuberculosis* specific antigenic and immunogenic properties. Other embodiments are compositions containing the polypeptide, or functional segment, and a pharmaceutically acceptable diluent or filler. Compositions of the invention can also contain at least two (e. g., three, four, five, six, seven, eight, nine, ten, twelve, fifteen, or twenty) polypeptides of the *Mycobacterium tuberculosis* complex, or functional segments thereof, with at least one of the at least two (e. g., two, three, four, five, six, seven, or eight) polypeptides having the sequence of one of the above described polypeptides of the invention. The polypeptides will preferably be those not encoded by the genome of cells of the BCG strain of *M. bovis*.

The invention also features methods of diagnosis. One embodiment is a method involving: (a) administration of one of the above polypeptide compositions to a subject suspected of having or being susceptible to *Mycobacterium tuberculosis* infection; and (b) detecting an immune response in the subject to the composition, as an indication that the subject has or is susceptible to *Mycobacterium tuberculosis* infection. An example of such a method is a skin test in which the test substance (e. g., compositions containing one or more of MTBN1-MTBN8) is injected intradermally into the subject and in which a skin delayed-type hypersensitivity response is tested for. Another embodiment is a method that involves: (a) providing a population of cells containing CD4 T lymphocytes from a subject; (b) providing a population of cells containing antigen presenting cells (APC) expressing a major histocompatibility complex (MHC) class II molecule expressed by the subject; (c) contacting the CD4 lymphocytes of (a) with the APC of (b) in the presence of one or more of the polypeptides, functional segments, and or polypeptide compositions of the invention; and (d) determining the ability of the CD4 lymphocytes to respond to the polypeptide, as an indication that the subject has or is susceptible to *Mycobacterium tuberculosis* infection. Another diagnostic method of the invention involves: (a) contacting a polypeptide, a functional segment, or a polypeptide/functional segment composition of the invention with a bodily fluid of a subject; (b) detecting the presence of binding of antibody to the polypeptide, functional segment, or polypeptide/functional segment composition, as an indication that the subject has or is susceptible to *Mycobacterium tuberculosis* infection.

Also encompassed by the invention are methods of vaccination. These methods involve administration of any of the above polypeptides, functional segments, or DNA compositions to a subject. The compositions can be administered alone or with one or more of the other compositions.

As used herein, an "isolated DNA molecule" is a DNA which is one or both of: not immediately contiguous with one or both of the coding sequences with which it is immediately contiguous (i.e., one at the 5' end and one at the 3' end) in the naturally-occurring genome of the organism from which the DNA is derived; or which is substantially free of DNA sequence with which it occurs in the organism from which the DNA is derived. The term includes, for example, a recombinant DNA which incorporated into a vector, e.g., into an autonomously replicating plasmid or virus, or into the genomic DNA of a prokaryote or eukaryote, or which exists as a separate molecule (e.g., a cDNA or a genomic fragment produced by PCR or restriction endonuclease treatment) independent of other DNA sequences. Isolated DNA also includes a recombinant DNA which is part of a hybrid DNA encoding additional *M. tuberculosis* polypeptide sequences.

"DNA molecules" include cDNA, genomic DNA, and synthetic (e.g., chemically synthesized) DNA. Where single-stranded, the DNA molecule may be a sense strand or an antisense strand.

An "isolated polypeptide" of the invention is a polypeptide which either has no naturally-occurring counterpart, or has been separated or purified from components which naturally accompany it, e.g., in *M. tuberculosis* bacteria.

Typically, the polypeptide is considered "isolated" when it is at least 70%, by dry weight, free from the proteins and naturally-occurring organic molecules with which it is naturally associated.

Preferably, a preparation of a polypeptide of the invention is at least 80%, more preferably at least 90%, and most preferably at least 99%, by dry weight, the peptide of the invention. Since a polypeptide that is chemically synthesized is, by its nature, separated from the components that naturally accompany it, the synthetic polypeptide is "isolated."

An isolated polypeptide of the invention can be obtained, for example, by extraction from a natural source (e.g., *M. tuberculosis* bacteria); by expression of a recombinant nucleic acid encoding the polypeptide; or by chemical synthesis. A polypeptide that is produced in a cellular system different from the source from which it naturally originates is "isolated," because it will be separated from components which naturally accompany it. The extent of isolation or purity can be measured by any appropriate method, e.g., column chromatography, polyacrylamide gel electrophoresis, or HPLC analysis.

The polypeptides may contain a primary amino acid sequence that has been modified from those disclosed herein. Preferably these modifications consist of conservative amino acid substitutions. Conservative substitutions typically include substitutions within the following groups: glycine and alanine; valine, isoleucine, and leucine; aspartic acid and glutamic acid; asparagine and glutamine; serine and threonine; lysine and arginine; and phenylalanine and tyrosine.

The terms "protein" and "polypeptide" are used herein to describe any chain of amino acids, regardless of length or post-translational modification (for example, glycosylation or phosphorylation). Thus, the term "*Mycobacterium tuberculosis* polypeptide" includes full-length, naturally occurring *Mycobacterium tuberculosis* protein, as well a recombinantly or synthetically produced polypeptide that corresponds to a full-length naturally occurring *Mycobacterium tuberculosis* protein or to particular domains or portions of a naturally occurring protein. The term also encompasses a mature *Mycobacterium tuberculosis* polypeptide which has an added amino-terminal methionine (useful for expression in prokaryotic cells) or any short amino acid sequences useful for protein purification by affinity chromatography, e.g., polyhistidine for purification by metal chelate chromatography.

As used herein, "immunogenic" means capable of activating a primary or memory immune response. Immune responses include responses of CD4+ and CD8+ T lymphocytes and B-lymphocytes. In the case of T lymphocytes, such responses can be proliferative, and/or cytokine (e. g., interleukin (IL)-2, IL-3, IL-4, IL-5, IL-6, IL-12, IL-13, IL-15, tumor necrosis factor- $\alpha$  (TNF- $\alpha$ ), or interferon- $\gamma$  (IFN- $\gamma$ ))-producing, or they can result in generation of cytotoxic T-lymphocytes (CTL). B-lymphocyte responses can be those resulting in antibody production by the responding B lymphocytes.

As used herein, "antigenic" means capable of being recognized by either antibody molecules or antigen-specific

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T cell receptors (TCR) on activated effector T cells (e. g., cytokine-producing T cells or CTL).

Thus, polypeptides that have “*Mycobacterium tuberculosis* specific antigenic properties” are polypeptides that: (a) can be recognized by and bind to antibodies elicited in response to *Mycobacterium tuberculosis* organisms or wild-type *Mycobacterium tuberculosis* molecules (e. g., polypeptides); or (b) contain subsequences which, subsequent to processing of the polypeptide by appropriate antigen presenting cells (APC) and bound to appropriate major histocompatibility complex (MHC) molecules, are recognized by and bind to TCR on effector T cells elicited in response to *Mycobacterium tuberculosis* organisms or wild-type *Mycobacterium tuberculosis* molecules (e. g., polypeptides).

As used herein, polypeptides that have “*Mycobacterium tuberculosis* specific immunogenic properties” are polypeptides that: (a) can elicit the production of antibodies that recognize and bind to *Mycobacterium tuberculosis* organisms or wild-type *Mycobacterium tuberculosis* molecules (e. g., polypeptides); or (b) contain subsequences which, subsequent to processing of the polypeptide by appropriate antigen presenting cells (APC) and bound to appropriate major histocompatibility complex (MHC) molecules on the surface of the APC, activate T cells with TCR that recognize and bind to peptide fragments derived by processing by APC of *Mycobacterium tuberculosis* organisms or wild-type *Mycobacterium tuberculosis* molecules (e. g., polypeptides) and bound to MHC molecules on the surface of the APC. The immune responses elicited in response to the immunogenic polypeptides are preferably protective. As used herein, “protective” means preventing establishment of an infection or onset of a disease or lessening the severity of a disease existing in a subject. “Preventing” can include delaying onset, as well as partially or completely blocking progress of the disease.

As used herein, a “functional segment of a *Mycobacterium tuberculosis* polypeptide” is a segment of the polypeptide that has *Mycobacterium tuberculosis* specific antigenic and immunogenic properties.

Where a polypeptide, functional segment of a polypeptide, or a mixture of polypeptides and/or functional segments have been administered (e.g., by intradermal injection) to a subject for the purpose of testing for a *M. tuberculosis* infection or susceptibility to such an infection, “detecting an immune response” means examining the subject for signs of an immunological reaction to the administered material, e.g., reddening or swelling of the skin at the site of an intradermal injection. Where the subject has antibodies to the administered material, the response will generally be rapid, e.g., 1 minute to 24 hours. On the other hand, a memory or activated T cell reaction of pre-immunized T lymphocytes in the subject is generally slower, appearing only after 24 hours and being maximal at 24-96 hours.

As used herein, a “subject” can be a human subject or a non-human mammal such as a non-human primate, a horse, a bovine animal, a pig, a sheep, a goat, a dog, a cat, a rabbit, a guinea pig, a hamster, a rat, or a mouse.

Unless otherwise defined, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this

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invention pertains. In case of conflict, the present document, including definitions, will control. Preferred methods and materials are described below, although methods and materials similar or equivalent to those described herein can be used in the practice or testing of the present invention. Unless otherwise indicated, these materials and methods are illustrative only and are not intended to be limiting.

All publications, patent applications, patents and other references mentioned herein are illustrative only and not intended to be limiting.

Other features and advantages of the invention, e. g., methods of diagnosing *M. tuberculosis* infection, will be apparent from the following description, from the drawings and from the claims.

## BRIEF DESCRIPTION OF THE DRAWINGS

FIGS. 1A and 1B are a depiction of the amino acid sequences of *M. tuberculosis* polypeptides MTBN1-MTBN8 (SEQ ID NOS:1-8, respectively).

FIGS. 2A-2E are a depiction of the nucleotide sequences of the coding regions (mtbn1-mtbn8) encoding MTBN1-MTBN8 (SEQ ID NOS:9-16, respectively).

FIG. 3 is a bar graph showing the delayed-type hypersensitivity responses induced by intradermal injection of 3 different test reagents in female guinea pigs that had been either infected with *M. tuberculosis* cells or sensitized with BCG or *M. avium* cells.

## DETAILED DESCRIPTION

The genome of *M. tuberculosis* [Cole et al. (1998) Nature 393: 537-544] contains open reading frames (ORFs) that have been deleted from the avirulent BCG strain.

The polypeptides encoded by these ORFs are designated herein “*M. tuberculosis* BCG Negative” polypeptides (“MTBN”) and the ORFs are designated “mtbn.” The invention is based on the discovery that a MTBN polypeptide (MTBN4) elicited a skin response in animals infected with *M. tuberculosis*, but not in animals sensitized to either BCG or *M. avium*, a non-*M. tuberculosis*-complex strain of mycobacteria (see Example 1 below). These findings indicate that MTBN (e.g., MTBN1-MTBN8) can be used in diagnostic tests that discriminate infection of a subject by *M. tuberculosis* from exposure to both mycobacteria other than the *M. tuberculosis*-complex and BCG. The *M. tuberculosis*-complex includes *M. tuberculosis*, *M. bovis*, *M. microti*, and *M. africanum*. Thus they can be used to discriminate subjects exposed to *M. tuberculosis*, and thus potentially having or being in danger of having tuberculosis, from subjects that have been vaccinated with BCG, the most widely used tuberculosis vaccine. Diagnostic assays that are capable of such discrimination represent a major advance that will greatly reduce wasted effort and consequent costs resulting from further diagnostic tests and/or therapeutic procedures in subjects that have given positive results in less discriminatory diagnostic tests.

Furthermore, the results in Example 1 show that MTBN4, as expressed by whole viable *M. tuberculosis* organisms, is capable of inducing a strong immune response in subjects infected with the organisms and thus has the potential to be a vaccine.

The MTBN polypeptides of the invention include, for example, polypeptides encoded within the RD1, RD2, and RD3 regions of the *M. tuberculosis* genome [Mahairas et al. (1996) J. Bacteriol. 178: 1274-1282]. Of particular interest are polypeptides encoded by ORFs within the RD1 region of the *M. tuberculosis* genome. However, the invention is not restricted to the RD1, RD2, and RD3 region encoded polypeptides and includes any polypeptides encoded by ORFs contained in the genome of one or more members of the *M. tuberculosis* genome and not contained in the genome of BCG. The amino acid sequences of MTBN1-MTBN8 are shown in FIGS. 1A and 1B and the nucleotide sequences of mtbn1-mtbn8 are shown in FIGS. 2A-2E.

The invention encompasses: (a) isolated DNA molecules containing mtbn sequences (e.g., mtbn1-mtbn8) encoding MTBN polypeptides (e.g., MTBN1-MTBN8) and isolated portions of such DNA molecules that encode polypeptide segments having antigenic and immunogenic properties (i.e., functional segments); (b) the MTBN polypeptides themselves (e.g., MTBN1-MTBN8) and functional segments of them; (c) antibodies (including antigen binding fragments, e.g., F (ab')<sub>2</sub>, Fab, Fv, and single chain Fv fragments of such antibodies) that bind to the MTBN polypeptides (e.g., MTBN1-MTBN8) and functional segments; (d) nucleic acid molecules (e.g., vectors) containing and capable of expressing one or more of the mtbn (e.g., mtbn1-mtbn8) sequences and portions of DNA molecules; (e) cells (e.g., bacterial, yeast, insect, or mammalian cells) transformed by such vectors; (f) compositions containing vectors encoding one or more *M. tuberculosis* polypeptides (or functional segments) including both the MTBN (e.g., MTBN1-MTBN8) polypeptides (or functional segments thereof) and previously described *M. tuberculosis* polypeptides such as ESAT-6, 14 kDa antigen, MPT63, 19 kDa antigen, MPT64, MPT51, MTC28, 38 kDa antigen, 45/47 kDa antigen, MPB70, Ag85 complex, MPT53, and KatG (see also U.S. Pat. No. 6,087,163); (g) compositions containing one or more *M. tuberculosis* polypeptides (or functional segments), including both the polypeptides of the invention and previously described *M. tuberculosis* polypeptides such as those described above; (h) compositions containing one or more of the antibodies described in (c); (i) methods of diagnosis involving either (1) administration (e.g., intradermal injection) of any of the above polypeptide compositions to a subject suspected of having or being susceptible to *M. tuberculosis* infection, (2) in vitro testing of lymphocytes (B-lymphocytes, CD4 T lymphocytes, and CD8 T lymphocytes) from such a subject for responsiveness (e.g., by measuring cell proliferation, antibody production, cytokine production, or CTL activity) to any of the above polypeptide compositions, (3) testing of a bodily fluid (e.g., blood, saliva, plasma, serum, urine, or semen or a lavage such as a bronchoalveolar lavage, a vaginal lavage, or lower gastrointestinal lavage) for antibodies to the MTBN polypeptides (e.g., MTBN1-MTBN8) or functional segments thereof, or the above-described polypeptide compositions; (4) testing of a bodily fluid (e.g., as above) for the presence of *M. tuberculosis*, MTBN (e.g., MTBN1-MTBN8) polypeptides or functional segments thereof, or the above-described polypeptide compositions in assays using the antibodies described in (c); and (5) testing of a tissue (e.g., lung

or bronchial tissue) or a body fluid (e.g., as above) for the presence of nucleic acid molecules (e.g., DNA or RNA) encoding MTBN polypeptides (e.g., MTBN1-MTBN8) (or portions of such a nucleic acid molecules) using nucleic acid probes or primers having nucleotide sequences of the nucleic molecules, portions of the nucleic molecules, or the complements of such molecules; and (j) methods of vaccination involving administration to a subject of the compositions of either (f), (g), (h) or a combination of any two or even all 3 compositions.

With respect to diagnosis, purified MTBN proteins, functional segments of such proteins, or mixtures of proteins and/or the functional fragments have the above-described advantages of discriminating infection by *M. tuberculosis* from either infection by other bacteria, and in particular, non-pathogenic mycobacteria, or from exposure (by, for example, vaccination) to BCG.

Furthermore, compositions containing the proteins, functional segments of the proteins, or mixtures of the proteins and/or the functional segments allows for improved quality control since "batch-to-batch" variability is greatly reduced in comparison to complex mixtures such as purified protein derivative (PPD) of tuberculin.

The use of the above-described polypeptide and nucleic acid reagents for vaccination also provides for highly specific and effective immunization. Since the virulent *M. tuberculosis* polypeptides encoded by genes absent from avirulent BCG are likely to be mediators of virulence, immunity directed to them can be especially potent in terms of protective capacity. Where vaccination is performed with nucleic acids both in vivo and ex vivo methods can be used. In vivo methods involve administration of the nucleic acids themselves to the subject and ex vivo methods involve obtaining cells (e.g., bone marrow cells or fibroblasts) from the subject, transducing the cells with the nucleic acids, preferably selecting or enriching for successfully transduced cells, and administering the transduced cells to the subject. Alternatively, the cells that are transduced and administered to the subject can be derived from another subject. Methods of vaccination and diagnosis are described in greater detail in U.S. Pat. No. 6,087,163, the disclosure of which is incorporated herein by reference in its entirety.

The following example is meant to illustrate, not limit the invention.

#### Example 1

##### MTBN4 Elicits a Specific Skin Reaction in Guinea Pigs Infected with *M. tuberculosis*

Four groups of outbred female guinea pigs (18 per group) were used to test the usefulness of the MTBN4 polypeptide as a *M. tuberculosis*-specific diagnostic reagent. The four groups were treated as follows.

Group 1 animals were infected by aerosol with approximately 100 *M. tuberculosis* strain H37Rv cells.

Group 2 animals were sensitized intradermally with 106 live *M. bovis* BCG Japanese cells.

Group 3 animals were sensitized intradermally with 106 live *M. avium* cells.

Group 4 animals were mock-sensitized by intradermal injection with saline.

Seven weeks after infection or sensitization, the animals were injected intradermally with 1 µg of PPD (6 animals from each group), 2 µg of purified recombinant MPT64 (6 animals from each group), or 2 µg of MTBN4 (6 animals from each group). The diameter of the resulting erythema was measured 24 hours later. Data are expressed as mean diameter of erythema (in mm) and standard deviations are indicated (FIG. 3).

No erythema was detected in the group 4 animals with any test substance and thus no data are shown for this group. On the other hand, group 1 animals (solid bars) showed a significant response with all three test substances. Group 2 animals (open bars) showed a significant response to PPD and MPT64 but not MTBN4.

Group 3 animals showed a significant response to PPD only (hatched bars).

Thus, PPD which contains antigenic/immunogenic molecules common to the *M. tuberculosis*-complex as well as

other mycobacterial strains, gave the least discriminatory results in that it induced responses in animals infected with or sensitized to mycobacteria of the *M. tuberculosis*-complex (*M. tuberculosis* and BCG) as well as another non-pathogenic mycobacterium (*M. avium*).

While MPT64, which is encoded and expressed by both *M. tuberculosis* and BCG, did not elicit a response in animals infected with *M. avium*, it did elicit responses in both the *M. tuberculosis* infected and the BCG sensitized animals. Finally, MTBN4 elicited a response in only the *M. tuberculosis* animals. Thus, it induced the most specific response and, most importantly, allowed for discrimination between animals infected with *M. tuberculosis* and those sensitized to BCG.

Although the invention has been described with reference to the presently preferred embodiment, it should be understood that various modifications can be made without departing from the spirit of the invention. Accordingly, the invention is limited only by the following claims.

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#### SEQUENCE LISTING

<160> NUMBER OF SEQ ID NOS: 16

<210> SEQ ID NO 1

<211> LENGTH: 591

<212> TYPE: PRT

<213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 1

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Met Thr Ala Glu Pro Glu Val Arg Thr Leu Arg Glu Val Val Leu Asp
 1             5             10             15

Gln Leu Gly Thr Ala Glu Ser Arg Ala Tyr Lys Met Trp Leu Pro Pro
          20             25             30

Leu Thr Asn Pro Val Pro Leu Asn Glu Leu Ile Ala Arg Asp Arg Arg
 35             40             45

Gln Pro Leu Arg Phe Ala Leu Gly Ile Met Asp Glu Pro Arg Arg His
 50             55             60

Leu Gln Asp Val Trp Gly Val Asp Val Ser Gly Ala Gly Gly Asn Ile
 65             70             75             80

Gly Ile Gly Gly Ala Pro Gln Thr Gly Lys Ser Thr Leu Leu Gln Thr
          85             90             95

Met Val Met Ser Ala Ala Ala Thr His Ser Pro Arg Asn Val Gln Phe
          100             105             110

Tyr Cys Ile Asp Leu Gly Gly Gly Gly Leu Ile Tyr Leu Glu Asn Leu
          115             120             125

Pro His Val Gly Gly Val Ala Asn Arg Ser Glu Pro Asp Lys Val Asn
          130             135             140

Arg Val Val Ala Glu Met Gln Ala Val Met Arg Gln Arg Glu Thr Thr
          145             150             155             160

Phe Lys Glu His Arg Val Gly Ser Ile Gly Met Tyr Arg Gln Leu Arg
          165             170             175

Asp Asp Pro Ser Gln Pro Val Ala Ser Asp Pro Tyr Gly Asp Val Phe
          180             185             190

Leu Ile Ile Asp Gly Trp Pro Gly Phe Val Gly Glu Phe Pro Asp Leu
          195             200             205

Glu Gly Gln Val Gln Asp Leu Ala Ala Gln Gly Leu Ala Phe Gly Val
          210             215             220

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His Val Ile Ile Ser Thr Pro Arg Trp Thr Glu Leu Lys Ser Arg Val
225                230                235                240

Arg Asp Tyr Leu Gly Thr Lys Ile Glu Phe Arg Leu Gly Asp Val Asn
                245                250                255

Glu Thr Gln Ile Asp Arg Ile Thr Arg Glu Ile Pro Ala Asn Arg Pro
                260                265                270

Gly Arg Ala Val Ser Met Glu Lys His His Leu Met Ile Gly Val Pro
                275                280                285

Arg Phe Asp Gly Val His Ser Ala Asp Asn Leu Val Glu Ala Ile Thr
290                295                300

Ala Gly Val Thr Gln Ile Ala Ser Gln His Thr Glu Gln Ala Pro Pro
305                310                315                320

Val Arg Val Leu Pro Glu Arg Ile His Leu His Glu Leu Asp Pro Asn
                325                330                335

Pro Pro Gly Pro Glu Ser Asp Tyr Arg Thr Arg Trp Glu Ile Pro Ile
                340                345                350

Gly Leu Arg Glu Thr Asp Leu Thr Pro Ala His Cys His Met His Thr
355                360                365

Asn Pro His Leu Leu Ile Phe Gly Ala Ala Lys Ser Gly Lys Thr Thr
370                375                380

Ile Ala His Ala Ile Ala Arg Ala Ile Cys Ala Arg Asn Ser Pro Gln
385                390                395                400

Gln Val Arg Phe Met Leu Ala Asp Tyr Arg Ser Gly Leu Leu Asp Ala
                405                410                415

Val Pro Asp Thr His Leu Leu Gly Ala Gly Ala Ile Asn Arg Asn Ser
420                425                430

Ala Ser Leu Asp Glu Ala Val Gln Ala Leu Ala Val Asn Leu Lys Lys
435                440                445

Arg Leu Pro Pro Thr Asp Leu Thr Thr Ala Gln Leu Arg Ser Arg Ser
450                455                460

Trp Trp Ser Gly Phe Asp Val Val Leu Leu Val Asp Asp Trp His Met
465                470                475                480

Ile Val Gly Ala Ala Gly Gly Met Pro Pro Met Ala Pro Leu Ala Pro
                485                490                495

Leu Leu Pro Ala Ala Ala Asp Ile Gly Leu His Ile Ile Val Thr Cys
500                505                510

Gln Met Ser Gln Ala Tyr Lys Ala Thr Met Asp Lys Phe Val Gly Ala
515                520                525

Ala Phe Gly Ser Gly Ala Pro Thr Met Phe Leu Ser Gly Glu Lys Gln
530                535                540

Glu Phe Pro Ser Ser Glu Phe Lys Val Lys Arg Arg Pro Pro Gly Gln
545                550                555                560

Ala Phe Leu Val Ser Pro Asp Gly Lys Glu Val Ile Gln Ala Pro Tyr
565                570                575

Ile Glu Pro Pro Glu Glu Val Phe Ala Ala Pro Pro Ser Ala Gly
580                585                590

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&lt;210&gt; SEQ ID NO 2

&lt;211&gt; LENGTH: 99

&lt;212&gt; TYPE: PRT

&lt;213&gt; ORGANISM: Mycobacterium tuberculosis

&lt;400&gt; SEQUENCE: 2

Met Glu Lys Met Ser His Asp Pro Ile Ala Ala Asp Ile Gly Thr Gln

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1	5	10	15
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Val Ser Asp Asn Ala Leu His Gly Val Thr Ala Gly Ser Thr Ala Leu  
                   20                  25                  30

Thr Ser Val Thr Gly Leu Val Pro Ala Gly Ala Asp Glu Val Ser Ala  
                   35                  40                  45

Gln Ala Ala Thr Ala Phe Thr Ser Glu Gly Ile Gln Leu Leu Ala Ser  
                   50                  55                  60

Asn Ala Ser Ala Gln Asp Gln Leu His Arg Ala Gly Glu Ala Val Gln  
                   65                  70                  75                  80

Asp Val Ala Arg Thr Tyr Ser Gln Ile Asp Asp Gly Ala Ala Gly Val  
                   85                  90                  95

Phe Ala Glu

<210> SEQ ID NO 3  
 <211> LENGTH: 368  
 <212> TYPE: PRT  
 <213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 3

Met Leu Trp His Ala Met Pro Pro Glu Leu Asn Thr Ala Arg Leu Met	
1                  5                  10                  15	
Ala Gly Ala Gly Pro Ala Pro Met Leu Ala Ala Ala Ala Gly Trp Gln	
20                  25                  30	
Thr Leu Ser Ala Ala Leu Asp Ala Gln Ala Val Glu Leu Thr Ala Arg	
35                  40                  45	
Leu Asn Ser Leu Gly Glu Ala Trp Thr Gly Gly Gly Ser Asp Lys Ala	
50                  55                  60	
Leu Ala Ala Ala Thr Pro Met Val Val Trp Leu Gln Thr Ala Ser Thr	
65                  70                  75                  80	
Gln Ala Lys Thr Arg Ala Met Gln Ala Thr Ala Gln Ala Ala Ala Tyr	
85                  90                  95	
Thr Gln Ala Met Ala Thr Thr Pro Ser Leu Pro Glu Ile Ala Ala Asn	
100                  105                  110	
His Ile Thr Gln Ala Val Leu Thr Ala Thr Asn Phe Phe Gly Ile Asn	
115                  120                  125	
Thr Ile Pro Ile Ala Leu Thr Glu Met Asp Tyr Phe Ile Arg Met Trp	
130                  135                  140	
Asn Gln Ala Ala Leu Ala Met Glu Val Tyr Gln Ala Glu Thr Ala Val	
145                  150                  155                  160	
Asn Thr Leu Phe Glu Lys Leu Glu Pro Met Ala Ser Ile Leu Asp Pro	
165                  170                  175	
Gly Ala Ser Gln Ser Thr Thr Asn Pro Ile Phe Gly Met Pro Ser Pro	
180                  185                  190	
Gly Ser Ser Thr Pro Val Gly Gln Leu Pro Pro Ala Ala Thr Gln Thr	
195                  200                  205	
Leu Gly Gln Leu Gly Glu Met Ser Gly Pro Met Gln Gln Leu Thr Gln	
210                  215                  220	
Pro Leu Gln Gln Val Thr Ser Leu Phe Ser Gln Val Gly Gly Thr Gly	
225                  230                  235                  240	
Gly Gly Asn Pro Ala Asp Glu Glu Ala Ala Gln Met Gly Leu Leu Gly	
245                  250                  255	
Thr Ser Pro Leu Ser Asn His Pro Leu Ala Gly Gly Ser Gly Pro Ser	
260                  265                  270	

Ala Gly Ala Gly Leu Leu Arg Ala Glu Ser Leu Pro Gly Ala Gly Gly

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275	280	285
Ser Leu Thr Arg Thr Pro	Leu Met Ser Gln Leu	Ile Glu Lys Pro Val
290	295	300
Ala Pro Ser Val Met Pro	Ala Ala Ala Ala Gly	Ser Ser Ala Thr Gly
305	310	315
Gly Ala Ala Pro Val Gly	Ala Gly Ala Met Gly	Gln Gly Ala Gln Ser
325	330	335
Gly Gly Ser Thr Arg Pro	Gly Leu Val Ala Pro	Ala Pro Leu Ala Gln
340	345	350
Glu Arg Glu Glu Asp Asp	Glu Asp Asp Trp Asp	Glu Glu Asp Asp Trp
355	360	365

<210> SEQ ID NO 4  
 <211> LENGTH: 100  
 <212> TYPE: PRT  
 <213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 4

Met Ala Glu Met Lys Thr	Asp Ala Ala Thr	Leu Ala Gln Glu Ala Gly
1	5	10
Asn Phe Glu Arg Ile Ser	Gly Asp Leu Lys Thr	Gln Ile Asp Gln Val
20	25	30
Glu Ser Thr Ala Gly Ser	Leu Gln Gly Gln Trp	Arg Gly Ala Ala Gly
35	40	45
Thr Ala Ala Gln Ala Ala	Val Val Arg Phe Gln	Glu Ala Ala Asn Lys
50	55	60
Gln Lys Gln Glu Leu Asp	Glu Ile Ser Thr Asn	Ile Arg Gln Ala Gly
65	70	75
Val Gln Tyr Ser Arg Ala	Asp Glu Glu Gln Gln	Gln Ala Leu Ser Ser
85	90	95

Gln Met Gly Phe  
100

<210> SEQ ID NO 5  
 <211> LENGTH: 666  
 <212> TYPE: PRT  
 <213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 5

Met Ala Ala Asp Tyr Asp	Lys Leu Phe Arg	Pro His Glu Gly Met Glu
1	5	10
Ala Pro Asp Asp Met Ala	Ala Gln Pro Phe Phe	Asp Pro Ser Ala Ser
20	25	30
Phe Pro Pro Ala Pro Ala	Ser Ala Asn Leu Pro	Lys Pro Asn Gly Gln
35	40	45
Thr Pro Pro Pro Thr Ser	Asp Asp Leu Ser Glu	Arg Phe Val Ser Ala
50	55	60
Pro Pro Pro Pro Pro Pro	Pro Pro Pro Pro Pro	Pro Pro Thr Pro Met
65	70	75
Pro Ile Ala Ala Gly Glu	Pro Pro Ser Pro Glu	Pro Ala Ala Ser Lys
85	90	95
Pro Pro Thr Pro Pro Met	Pro Ile Ala Gly Pro	Glu Pro Ala Pro Pro
100	105	110
Lys Pro Pro Thr Pro Pro	Met Pro Ile Ala Gly	Pro Glu Pro Ala Pro
115	120	125
Pro Lys Pro Pro Thr Pro	Pro Met Pro Ile Ala	Gly Pro Ala Pro Thr

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130	135	140
Pro Thr Glu Ser Gln Leu Ala Pro Pro Arg Pro Pro Thr Pro Gln Thr		
145	150	155
Pro Thr Gly Ala Pro Gln Gln Pro Glu Ser Pro Ala Pro His Val Pro		
	165	170
		175
Ser His Gly Pro His Gln Pro Arg Arg Thr Ala Pro Ala Pro Pro Trp		
	180	185
		190
Ala Lys Met Pro Ile Gly Glu Pro Pro Pro Ala Pro Ser Arg Pro Ser		
	195	200
		205
Ala Ser Pro Ala Glu Pro Pro Thr Arg Pro Ala Pro Gln His Ser Arg		
	210	215
		220
Arg Ala Arg Arg Gly His Arg Tyr Arg Thr Asp Thr Glu Arg Asn Val		
	225	230
		235
		240
Gly Lys Val Ala Thr Gly Pro Ser Ile Gln Ala Arg Leu Arg Ala Glu		
	245	250
		255
Glu Ala Ser Gly Ala Gln Leu Ala Pro Gly Thr Glu Pro Ser Pro Ala		
	260	265
		270
Pro Leu Gly Gln Pro Arg Ser Tyr Leu Ala Pro Pro Thr Arg Pro Ala		
	275	280
		285
Pro Thr Glu Pro Pro Pro Ser Pro Ser Pro Gln Arg Asn Ser Gly Arg		
	290	295
		300
Arg Ala Glu Arg Arg Val His Pro Asp Leu Ala Ala Gln His Ala Ala		
	305	310
		315
		320
Ala Gln Pro Asp Ser Ile Thr Ala Ala Thr Thr Gly Gly Arg Arg Arg		
	325	330
		335
Lys Arg Ala Ala Pro Asp Leu Asp Ala Thr Gln Lys Ser Leu Arg Pro		
	340	345
		350
Ala Ala Lys Gly Pro Lys Val Lys Lys Val Lys Pro Gln Lys Pro Lys		
	355	360
		365
Ala Thr Lys Pro Pro Lys Val Val Ser Gln Arg Gly Trp Arg His Trp		
	370	375
		380
Val His Ala Leu Thr Arg Ile Asn Leu Gly Leu Ser Pro Asp Glu Lys		
	385	390
		395
		400
Tyr Glu Leu Asp Leu His Ala Arg Val Arg Arg Asn Pro Arg Gly Ser		
	405	410
		415
Tyr Gln Ile Ala Val Val Gly Leu Lys Gly Gly Ala Gly Lys Thr Thr		
	420	425
		430
Leu Thr Ala Ala Leu Gly Ser Thr Leu Ala Gln Val Arg Ala Asp Arg		
	435	440
		445
Ile Leu Ala Leu Asp Ala Asp Pro Gly Ala Gly Asn Leu Ala Asp Arg		
	450	455
		460
Val Gly Arg Gln Ser Gly Ala Thr Ile Ala Asp Val Leu Ala Glu Lys		
	465	470
		475
		480
Glu Leu Ser His Tyr Asn Asp Ile Arg Ala His Thr Ser Val Asn Ala		
	485	490
		495
Val Asn Leu Glu Val Leu Pro Ala Pro Glu Tyr Ser Ser Ala Gln Arg		
	500	505
		510
Ala Leu Ser Asp Ala Asp Trp His Phe Ile Ala Asp Pro Ala Ser Arg		
	515	520
		525
Phe Tyr Asn Leu Val Leu Ala Asp Cys Gly Ala Gly Phe Phe Asp Pro		
	530	535
		540
Leu Thr Arg Gly Val Leu Ser Thr Val Ser Gly Val Val Val Val Ala		
	545	550
		555
		560

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Ser Val Ser Ile Asp Gly Ala Gln Gln Ala Ser Val Ala Leu Asp Trp  
565 570 575  
Leu Arg Asn Asn Gly Tyr Gln Asp Leu Ala Ser Arg Ala Cys Val Val  
580 585 590  
Ile Asn His Ile Met Pro Gly Glu Pro Asn Val Ala Val Lys Asp Leu  
595 600 605  
Val Arg His Phe Glu Gln Gln Val Gln Pro Gly Arg Val Val Val Met  
610 615 620  
Pro Trp Asp Arg His Ile Ala Ala Gly Thr Glu Ile Ser Leu Asp Leu  
625 630 635 640  
Leu Asp Pro Ile Tyr Lys Arg Lys Val Leu Glu Leu Ala Ala Ala Leu  
645 650 655  
Ser Asp Asp Phe Glu Arg Ala Gly Arg Arg  
660 665

<210> SEQ ID NO 6  
<211> LENGTH: 511  
<212> TYPE: PRT  
<213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 6

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Ala Ala Arg Pro Ala Thr Thr Arg Val Thr Ile Leu Thr Gly Arg Arg  
20 25 30  
Met Thr Asp Leu Val Leu Pro Ala Ala Val Pro Met Glu Thr Tyr Ile  
35 40 45  
Asp Asp Thr Val Ala Val Leu Ser Glu Val Leu Glu Asp Thr Pro Ala  
50 55 60  
Asp Val Leu Gly Gly Phe Asp Phe Thr Ala Gln Gly Val Trp Ala Phe  
65 70 75 80  
Ala Arg Pro Gly Ser Pro Pro Leu Lys Leu Asp Gln Ser Leu Asp Asp  
85 90 95  
Ala Gly Val Val Asp Gly Ser Leu Leu Thr Leu Val Ser Val Ser Arg  
100 105 110  
Thr Glu Arg Tyr Arg Pro Leu Val Glu Asp Val Ile Asp Ala Ile Ala  
115 120 125  
Val Leu Asp Glu Ser Pro Glu Phe Asp Arg Thr Ala Leu Asn Arg Phe  
130 135 140  
Val Gly Ala Ala Ile Pro Leu Leu Thr Ala Pro Val Ile Gly Met Ala  
145 150 155 160  
Met Arg Ala Trp Trp Glu Thr Gly Arg Ser Leu Trp Trp Pro Leu Ala  
165 170 175  
Ile Gly Ile Leu Gly Ile Ala Val Leu Val Gly Ser Phe Val Ala Asn  
180 185 190  
Arg Phe Tyr Gln Ser Gly His Leu Ala Glu Cys Leu Leu Val Thr Thr  
195 200 205  
Tyr Leu Leu Ile Ala Thr Ala Ala Ala Leu Ala Val Pro Leu Pro Arg  
210 215 220  
Gly Val Asn Ser Leu Gly Ala Pro Gln Val Ala Gly Ala Ala Thr Ala  
225 230 235 240  
Val Leu Phe Leu Thr Leu Met Thr Arg Gly Gly Pro Arg Lys Arg His  
245 250 255  
Glu Leu Ala Ser Phe Ala Val Ile Thr Ala Ile Ala Val Ile Ala Ala

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260	265	270
Ala Ala Ala Phe Gly Tyr Gly Tyr Gln Asp Trp Val Pro Ala Gly Gly		
275	280	285
Ile Ala Phe Gly Leu Phe Ile Val Thr Asn Ala Ala Lys Leu Thr Val		
290	295	300
Ala Val Ala Arg Ile Ala Leu Pro Pro Ile Pro Val Pro Gly Glu Thr		
305	310	315
Val Asp Asn Glu Glu Leu Leu Asp Pro Val Ala Thr Pro Glu Ala Thr		
325	330	335
Ser Glu Glu Thr Pro Thr Trp Gln Ala Ile Ile Ala Ser Val Pro Ala		
340	345	350
Ser Ala Val Arg Leu Thr Glu Arg Ser Lys Leu Ala Lys Gln Leu Leu		
355	360	365
Ile Gly Tyr Val Thr Ser Gly Thr Leu Ile Leu Ala Ala Gly Ala Ile		
370	375	380
Ala Val Val Val Arg Gly His Phe Phe Val His Ser Leu Val Val Ala		
385	390	395
Gly Leu Ile Thr Thr Val Cys Gly Phe Arg Ser Arg Leu Tyr Ala Glu		
405	410	415
Arg Trp Cys Ala Trp Ala Leu Leu Ala Ala Thr Val Ala Ile Pro Thr		
420	425	430
Gly Leu Thr Ala Lys Leu Ile Ile Trp Tyr Pro His Tyr Ala Trp Leu		
435	440	445
Leu Leu Ser Val Tyr Leu Thr Val Ala Leu Val Ala Leu Val Val Val		
450	455	460
Gly Ser Met Ala His Val Arg Arg Val Ser Pro Val Val Lys Arg Thr		
465	470	475
Leu Glu Leu Ile Asp Gly Ala Met Ile Ala Ala Ile Ile Pro Met Leu		
485	490	495
Leu Trp Ile Thr Gly Val Tyr Asp Thr Val Arg Asn Ile Arg Phe		
500	505	510

<210> SEQ ID NO 7  
 <211> LENGTH: 280  
 <212> TYPE: PRT  
 <213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 7

Met Ala Glu Pro Leu Ala Val Asp Pro Thr Gly Leu Ser Ala Ala Ala
1 5 10 15
Ala Lys Leu Ala Gly Leu Val Phe Pro Gln Pro Pro Ala Pro Ile Ala
20 25 30
Val Ser Gly Thr Asp Ser Val Val Ala Ala Ile Asn Glu Thr Met Pro
35 40 45
Ser Ile Glu Ser Leu Val Ser Asp Gly Leu Pro Gly Val Lys Ala Ala
50 55 60
Leu Thr Arg Thr Ala Ser Asn Met Asn Ala Ala Ala Asp Val Tyr Ala
65 70 75 80
Lys Thr Asp Gln Ser Leu Gly Thr Ser Leu Ser Gln Tyr Ala Phe Gly
85 90 95
Ser Ser Gly Glu Gly Leu Ala Gly Val Ala Ser Val Gly Gly Gln Pro
100 105 110
Ser Gln Ala Thr Gln Leu Leu Ser Thr Pro Val Ser Gln Val Thr Thr
115 120 125

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Gln Leu Gly Glu Thr Ala Ala Glu Leu Ala Pro Arg Val Val Ala Thr  
 130 135 140

Val Pro Gln Leu Val Gln Leu Ala Pro His Ala Val Gln Met Ser Gln  
 145 150 155 160

Asn Ala Ser Pro Ile Ala Gln Thr Ile Ser Gln Thr Ala Gln Gln Ala  
 165 170 175

Ala Gln Ser Ala Gln Gly Gly Ser Gly Pro Met Pro Ala Gln Leu Ala  
 180 185 190

Ser Ala Glu Lys Pro Ala Thr Glu Gln Ala Glu Pro Val His Glu Val  
 195 200 205

Thr Asn Asp Asp Gln Gly Asp Gln Gly Asp Val Gln Pro Ala Glu Val  
 210 215 220

Val Ala Ala Ala Arg Asp Glu Gly Ala Gly Ala Ser Pro Gly Gln Gln  
 225 230 235 240

Pro Gly Gly Gly Val Pro Ala Gln Ala Met Asp Thr Gly Ala Gly Ala  
 245 250 255

Arg Pro Ala Ala Ser Pro Leu Ala Ala Pro Val Asp Pro Ser Thr Pro  
 260 265 270

Ala Pro Ser Thr Thr Thr Thr Leu  
 275 280

<210> SEQ ID NO 8  
 <211> LENGTH: 729  
 <212> TYPE: PRT  
 <213> ORGANISM: Mycobacterium tuberculosis

<400> SEQUENCE: 8

Met Ser Ile Thr Arg Pro Thr Gly Ser Tyr Ala Arg Gln Met Leu Asp  
 1 5 10 15

Pro Gly Gly Trp Val Glu Ala Asp Glu Asp Thr Phe Tyr Asp Arg Ala  
 20 25 30

Gln Glu Tyr Ser Gln Val Leu Gln Arg Val Thr Asp Val Leu Asp Thr  
 35 40 45

Cys Arg Gln Gln Lys Gly His Val Phe Glu Gly Gly Leu Trp Ser Gly  
 50 55 60

Gly Ala Ala Asn Ala Ala Asn Gly Ala Leu Gly Ala Asn Ile Asn Gln  
 65 70 75 80

Leu Met Thr Leu Gln Asp Tyr Leu Ala Thr Val Ile Thr Trp His Arg  
 85 90 95

His Ile Ala Gly Leu Ile Glu Gln Ala Lys Ser Asp Ile Gly Asn Asn  
 100 105 110

Val Asp Gly Ala Gln Arg Glu Ile Asp Ile Leu Glu Asn Asp Pro Ser  
 115 120 125

Leu Asp Ala Asp Glu Arg His Thr Ala Ile Asn Ser Leu Val Thr Ala  
 130 135 140

Thr His Gly Ala Asn Val Ser Leu Val Ala Glu Thr Ala Glu Arg Val  
 145 150 155 160

Leu Glu Ser Lys Asn Trp Lys Pro Pro Lys Asn Ala Leu Glu Asp Leu  
 165 170 175

Leu Gln Gln Lys Ser Pro Pro Pro Pro Asp Val Pro Thr Leu Val Val  
 180 185 190

Pro Ser Pro Gly Thr Pro Gly Thr Pro Gly Thr Pro Ile Thr Pro Gly  
 195 200 205

Thr Pro Ile Thr Pro Gly Thr Pro Ile Thr Pro Ile Pro Gly Ala Pro  
 210 215 220

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Val Thr Pro Ile Thr Pro Thr Pro Gly Thr Pro Val Thr Pro Val Thr	225	230	235	240
Pro Gly Lys Pro Val Thr Pro Val Thr Pro Val Lys Pro Gly Thr Pro		245	250	255
Gly Glu Pro Thr Pro Ile Thr Pro Val Thr Pro Pro Val Ala Pro Ala		260	265	270
Thr Pro Ala Thr Pro Ala Thr Pro Val Thr Pro Ala Pro Ala Pro His		275	280	285
Pro Gln Pro Ala Pro Ala Pro Ala Pro Ser Pro Gly Pro Gln Pro Val	290	295	300	
Thr Pro Ala Thr Pro Gly Pro Ser Gly Pro Ala Thr Pro Gly Thr Pro	305	310	315	320
Gly Gly Glu Pro Ala Pro His Val Lys Pro Ala Ala Leu Ala Glu Gln		325	330	335
Pro Gly Val Pro Gly Gln His Ala Gly Gly Gly Thr Gln Ser Gly Pro		340	345	350
Ala His Ala Asp Glu Ser Ala Ala Ser Val Thr Pro Ala Ala Ala Ser		355	360	365
Gly Val Pro Gly Ala Arg Ala Ala Ala Ala Pro Ser Gly Thr Ala	370	375	380	
Val Gly Ala Gly Ala Arg Ser Ser Val Gly Thr Ala Ala Ala Ser Gly	385	390	395	400
Ala Gly Ser His Ala Ala Thr Gly Arg Ala Pro Val Ala Thr Ser Asp		405	410	415
Lys Ala Ala Ala Pro Ser Thr Arg Ala Ala Ser Ala Arg Thr Ala Pro		420	425	430
Pro Ala Arg Pro Pro Ser Thr Asp His Ile Asp Lys Pro Asp Arg Ser	435	440	445	
Glu Ser Ala Asp Asp Gly Thr Pro Val Ser Met Ile Pro Val Ser Ala	450	455	460	
Ala Arg Ala Ala Arg Asp Ala Ala Thr Ala Ala Ala Ser Ala Arg Gln	465	470	475	480
Arg Gly Arg Gly Asp Ala Leu Arg Leu Ala Arg Arg Ile Ala Ala Ala		485	490	495
Leu Asn Ala Ser Asp Asn Asn Ala Gly Asp Tyr Gly Phe Phe Trp Ile		500	505	510
Thr Ala Val Thr Thr Asp Gly Ser Ile Val Val Ala Asn Ser Tyr Gly	515	520	525	
Leu Ala Tyr Ile Pro Asp Gly Met Glu Leu Pro Asn Lys Val Tyr Leu	530	535	540	
Ala Ser Ala Asp His Ala Ile Pro Val Asp Glu Ile Ala Arg Cys Ala	545	550	555	560
Thr Tyr Pro Val Leu Ala Val Gln Ala Trp Ala Ala Phe His Asp Met		565	570	575
Thr Leu Arg Ala Val Ile Gly Thr Ala Glu Gln Leu Ala Ser Ser Asp		580	585	590
Pro Gly Val Ala Lys Ile Val Leu Glu Pro Asp Asp Ile Pro Glu Ser		595	600	605
Gly Lys Met Thr Gly Arg Ser Arg Leu Glu Val Val Asp Pro Ser Ala	610	615	620	
Ala Ala Gln Leu Ala Asp Thr Thr Asp Gln Arg Leu Leu Asp Leu Leu	625	630	635	640



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Pro Pro Ala Pro Val Asp Val Asn Pro Pro Gly Asp Glu Arg His Met  
645 650 655

Leu Trp Phe Glu Leu Met Lys Pro Met Thr Ser Thr Ala Thr Gly Arg  
660 665 670

Glu Ala Ala His Leu Arg Ala Phe Arg Ala Tyr Ala Ala His Ser Gln  
675 680 685

Glu Ile Ala Leu His Gln Ala His Thr Ala Thr Asp Ala Ala Val Gln  
690 695 700

Arg Val Ala Val Ala Asp Trp Leu Tyr Trp Gln Tyr Val Thr Gly Leu  
705 710 715 720

Leu Asp Arg Ala Leu Ala Ala Ala Cys  
725

<210> SEQ ID NO 9  
 <211> LENGTH: 1776  
 <212> TYPE: DNA  
 <213> ORGANISM: Mycobacterium tuberculosis  
 <220> FEATURE:  
 <221> NAME/KEY: CDS  
 <222> LOCATION: (1)...(1773)

<400> SEQUENCE: 9

atg act gct gaa ccg gaa gta ccg acg ctg cgc gag gtt gtg ctg gac	48
Met Thr Ala Glu Pro Glu Val Arg Thr Leu Arg Glu Val Val Leu Asp	
1 5 10 15	
cag ctc ggc act gct gaa tcg cgt gcg tac aag atg tgg ctg ccg ccg	96
Gln Leu Gly Thr Ala Glu Ser Arg Ala Tyr Lys Met Trp Leu Pro Pro	
20 25 30	
ttg acc aat ccg gtc ccg ctc aac gag ctc atc gcc cgt gat ccg cga	144
Leu Thr Asn Pro Val Pro Leu Asn Glu Leu Ile Ala Arg Asp Arg Arg	
35 40 45	
caa ccc ctg cga ttt gcc ctg ggg atc atg gat gaa ccg cgc cgc cat	192
Gln Pro Leu Arg Phe Ala Leu Gly Ile Met Asp Glu Pro Arg Arg His	
50 55 60	
cta cag gat gtg tgg ggc gta gac gtt tcc ggg gcc ggc ggc aac atc	240
Leu Gln Asp Val Trp Gly Val Asp Val Ser Gly Ala Gly Gly Asn Ile	
65 70 75 80	
ggg att ggg ggc gca cct caa acc ggg aag tcg acg cta ctg cag acg	288
Gly Ile Gly Gly Ala Pro Gln Thr Gly Lys Ser Thr Leu Leu Gln Thr	
85 90 95	
atg gtg atg tcg gcc gcc gcc aca cac tca ccg cgc aac gtt cag ttc	336
Met Val Met Ser Ala Ala Ala Thr His Ser Pro Arg Asn Val Gln Phe	
100 105 110	
tat tgc atc gac cta ggt ggc ggc ggg ctg atc tat ctc gaa aac ctt	384
Tyr Cys Ile Asp Leu Gly Gly Gly Gly Leu Ile Tyr Leu Glu Asn Leu	
115 120 125	
cca cac gtc ggt ggg gta gcc aat ccg tcc gag ccc gac aag gtc aac	432
Pro His Val Gly Gly Val Ala Asn Arg Ser Glu Pro Asp Lys Val Asn	
130 135 140	
cgg gtg gtc gca gag atg caa gcc gtc atg ccg caa ccg gaa acc acc	480
Arg Val Val Ala Glu Met Gln Ala Val Met Arg Gln Arg Glu Thr Thr	
145 150 155 160	
ttc aag gaa cac cga gtg ggc tcg atc ggg atg tac ccg cag ctg cgt	528
Phe Lys Glu His Arg Val Gly Ser Ile Gly Met Tyr Arg Gln Leu Arg	
165 170 175	
gac gat cca agt caa ccc gtt gcg tcc gat cca tac ggc gac gtc ttt	576
Asp Asp Pro Ser Gln Pro Val Ala Ser Asp Pro Tyr Gly Asp Val Phe	
180 185 190	
ctg atc atc gac gga tgg ccc ggt ttt gtc ggc gag ttc ccc gac ctt	624
Leu Ile Ile Asp Gly Trp Pro Gly Phe Val Gly Glu Phe Pro Asp Leu	

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195	200	205	
gag ggg cag gtt caa gat Glu Gly Gln Val Gln Asp 210	ctg gcc gcc cag ggg Leu Ala Ala Gln Gly 215	ctg gcg ttc ggc gtc Leu Ala Phe Gly Val 220	672
cac gtc atc atc tcc acg His Val Ile Ile Ser Thr 225	cca cgc tgg aca gag Pro Arg Trp Thr 230	ctg aag tcg cgt gtt Glu Leu Lys Ser Arg Val 235 240	720
cgc gac tac ctc ggc acc Arg Asp Tyr Leu Gly Thr 245	aag atc gag ttc cgg Lys Ile Glu Phe Arg 250	ctt ggt gac gtc aat Leu Gly Asp Val Asn 255	768
gaa acc cag atc gac cgg Glu Thr Gln Ile Asp Arg 260	att acc cgc gag atc ccg Ile Thr Arg Glu Ile Pro 265	gcg aat cgt ccg Ala Asn Arg Pro 270	816
ggg cgg gca gtg tgc atg Gly Arg Ala Val Ser Met 275	gaa aag cac cat ctg Glu Lys His His Leu Met 280	atg atc ggc gtg ccc Ile Gly Val Pro 285	864
agg ttc gac ggc gtg cac Arg Phe Asp Gly Val His 290	agc gcc gat aac ctg Ser Ala Asp Asn Leu 295	gtg gag gcg atc acc Val Glu Ala Ile Thr 300	912
gcg ggg gtg acg cag atc Ala Gly Val Thr Gln Ile 305	gct tcc cag cac acc Ile Ala Ser Gln His Thr 310	gaa cag gca cct ccg Glu Gln Ala Pro Pro 315 320	960
gtg cgg gtc ctg ccg gag Val Arg Val Leu Pro Glu 325	cgt atc cac ctg cac Arg Ile His Leu His 330	gaa ctc gac ccg aac Glu Leu Asp Pro Asn 335	1008
ccg ccg gga cca gag tcc Pro Pro Gly Pro Glu Ser 340	gac tac cgc act cgc Asp Tyr Arg Thr Arg 345	tgg gag att ccg atc Trp Glu Ile Pro Ile 350	1056
ggc ttg cgc gag acg gac Gly Leu Arg Glu Thr Asp 355	ctg acg ccg gct cac Leu Thr Pro Ala His 360	tgc cac atg cac acg Cys His Met His Thr 365	1104
aac ccg cac cta ctg atc Asn Pro His Leu Leu Ile 370	ttc ggt gcg gcc aaa Phe Gly Ala Ala Lys 375	tcg ggc aag acg acc Ser Gly Lys Thr Thr 380	1152
att gcc cac gcg atc gcg Ile Ala His Ala Ile Ala 385	gcg gcc att tgt gcc Arg Ala Ile Cys Ala 390	cga aac agt ccc cag Arg Asn Ser Pro Gln 395 400	1200
cag gtg cgg ttc atg ctc Gln Val Arg Phe Met 405	gcg gac tac cgc tcg Leu Ala Asp Tyr Arg 410	ggc ctg ctg gac gcg Gly Leu Leu Asp Ala 415	1248
gtg ccg gac acc cat ctg Val Pro Asp Thr His Leu 420	ctg ctg ggc gcc ggc Leu Leu Gly Ala Gly 425	gcg atc aac cgc aac agc Ala Ile Asn Arg Asn Ser 430	1296
gcg tcg cta gac gag gcc Ala Ser Leu Asp Glu Ala 435	gtt caa gca ctg gcg Val Gln Ala Leu Ala 440	gtc aac ctg aag aag Val Asn Leu Lys Lys 445	1344
cgg ttg ccg ccg acc gac Arg Leu Pro Pro Thr Asp 450	ctg acg acg gcg cag Leu Thr Thr Ala Gln 455	cta cgc tcg cgt tcg Leu Arg Ser Arg Ser 460	1392
tgg tgg agc gga ttt gac Trp Trp Ser Gly Phe Asp 465	gtc gtg ctt ctg gtc Val Val Leu Leu Val 470	gac gat tgg cac atg Asp Asp Trp His Met 475 480	1440
atc gtg ggt gcc gcc ggg Ile Val Gly Ala Ala Gly 485	ggg atg ccg ccg atg Gly Met Pro Pro Met 490	gca ccg ctg gcc ccg Ala Pro Leu Ala Pro 495	1488
tta ttg ccg gcg gcg gca Leu Leu Pro Ala Ala Ala 500	gat atc ggg ttg cac Asp Ile Gly Leu His 505	atc att gtc acc tgt Ile Ile Val Thr Cys 510	1536
cag atg agc cag gct tac aag gca acc atg gac aag ttc gtc ggc gcc			1584

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Gln Met Ser Gln Ala Tyr Lys Ala Thr Met Asp Lys Phe Val Gly Ala	
515 520 525	
gca ttc ggg tgc ggc gct ccg aca atg ttc ctt tgc ggc gag aag cag	1632
Ala Phe Gly Ser Gly Ala Pro Thr Met Phe Leu Ser Gly Glu Lys Gln	
530 535 540	
gaa ttc cca tcc agt gag ttc aag gtc aag cgg cgc ccc cct ggc cag	1680
Glu Phe Pro Ser Ser Glu Phe Lys Val Lys Arg Arg Pro Pro Gly Gln	
545 550 555 560	
gca ttt ctc gtc tgc cca gac ggc aaa gag gtc atc cag gcc ccc tac	1728
Ala Phe Leu Val Ser Pro Asp Gly Lys Glu Val Ile Gln Ala Pro Tyr	
565 570 575	
atc gag cct cca gaa gaa gtg ttc gca gca ccc cca agc gcc ggt	1773
Ile Glu Pro Pro Glu Val Phe Ala Ala Pro Pro Ser Ala Gly	
580 585 590	
taa	1776

<210> SEQ ID NO 10  
 <211> LENGTH: 300  
 <212> TYPE: DNA  
 <213> ORGANISM: Mycobacterium tuberculosis  
 <220> FEATURE:  
 <221> NAME/KEY: CDS  
 <222> LOCATION: (1)...(297)

<400> SEQUENCE: 10

atg gaa aaa atg tca cat gat ccg atc gct gcc gac att ggc acg caa	48
Met Glu Lys Met Ser His Asp Pro Ile Ala Ala Asp Ile Gly Thr Gln	
1 5 10 15	
gtg agc gac aac gct ctg cac ggc gtg acg gcc ggc tgc acg gcg ctg	96
Val Ser Asp Asn Ala Leu His Gly Val Thr Ala Gly Ser Thr Ala Leu	
20 25 30	
acg tgc gtg acc ggg ctg gtt ccc gcg ggg gcc gat gag gtc tcc gcc	144
Thr Ser Val Thr Gly Leu Val Pro Ala Gly Ala Asp Glu Val Ser Ala	
35 40 45	
caa gcg gcg acg gcg ttc aca tgc gag gcc atc caa ttg ctg gct tcc	192
Gln Ala Ala Thr Ala Phe Thr Ser Glu Gly Ile Gln Leu Leu Ala Ser	
50 55 60	
aat gca tgc gcc caa gac cag ctc cac cgt gcg ggc gaa gcg gtc cag	240
Asn Ala Ser Ala Gln Asp Gln Leu His Arg Ala Gly Glu Ala Val Gln	
65 70 75 80	
gac gtc gcc cgc acc tat tgc caa atc gac gac ggc gcc gcc ggc gtc	288
Asp Val Ala Arg Thr Tyr Ser Gln Ile Asp Asp Gly Ala Ala Gly Val	
85 90 95	
ttc gcc gaa tag	300
Phe Ala Glu	

<210> SEQ ID NO 11  
 <211> LENGTH: 1107  
 <212> TYPE: DNA  
 <213> ORGANISM: Mycobacterium tuberculosis  
 <220> FEATURE:  
 <221> NAME/KEY: CDS  
 <222> LOCATION: (1)...(1104)

<400> SEQUENCE: 11

atg ctg tgg cac gca atg cca ccg gag cta aat acc gca cgg ctg atg	48
Met Leu Trp His Ala Met Pro Pro Glu Leu Asn Thr Ala Arg Leu Met	
1 5 10 15	
gcc gcc gcg ggt ccg gct cca atg ctt gcg gcg gcc gcg gga tgg cag	96
Ala Gly Ala Gly Pro Ala Pro Met Leu Ala Ala Ala Ala Gly Trp Gln	
20 25 30	
acg ctt tgc gcg gct ctg gac gct cag gcc gtc gag ttg acc gcg cgc	144

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Thr	Leu	Ser	Ala	Ala	Leu	Asp	Ala	Gln	Ala	Val	Glu	Leu	Thr	Ala	Arg	
		35					40					45				
ctg	aac	tct	ctg	gga	gaa	gcc	tgg	act	gga	ggg	ggc	agc	gac	aag	gcg	192
Leu	Asn	Ser	Leu	Gly	Glu	Ala	Trp	Thr	Gly	Gly	Gly	Ser	Asp	Lys	Ala	
	50				55					60						
ctt	gcg	gct	gca	acg	ccg	atg	gtg	gtc	tgg	cta	caa	acc	gcg	tca	aca	240
Leu	Ala	Ala	Ala	Thr	Pro	Met	Val	Val	Trp	Leu	Gln	Thr	Ala	Ser	Thr	
	65				70				75					80		
cag	gcc	aag	acc	cgt	gcg	atg	cag	gcg	acg	gcg	caa	gcc	gcg	gca	tac	288
Gln	Ala	Lys	Thr	Arg	Ala	Met	Gln	Ala	Thr	Ala	Gln	Ala	Ala	Ala	Tyr	
				85				90						95		
acc	cag	gcc	atg	gcc	acg	acg	ccg	tcg	ctg	ccg	gag	atc	gcc	gcc	aac	336
Thr	Gln	Ala	Met	Ala	Thr	Thr	Pro	Ser	Leu	Pro	Glu	Ile	Ala	Ala	Asn	
			100				105						110			
cac	atc	acc	cag	gcc	gtc	ctt	acg	gcc	acc	aac	ttc	ttc	ggg	atc	aac	384
His	Ile	Thr	Gln	Ala	Val	Leu	Thr	Ala	Thr	Asn	Phe	Phe	Gly	Ile	Asn	
			115			120							125			
acg	atc	ccg	atc	gcg	ttg	acc	gag	atg	gat	tat	ttc	atc	cgt	atg	tgg	432
Thr	Ile	Pro	Ile	Ala	Leu	Thr	Glu	Met	Asp	Tyr	Phe	Ile	Arg	Met	Trp	
	130				135					140						
aac	cag	gca	gcc	ctg	gca	atg	gag	gtc	tac	cag	gcc	gag	acc	gcg	gtt	480
Asn	Gln	Ala	Ala	Leu	Ala	Met	Glu	Val	Tyr	Gln	Ala	Glu	Thr	Ala	Val	
	145				150					155					160	
aac	acg	ctt	ttc	gag	aag	ctc	gag	ccg	atg	gcg	tcg	atc	ctt	gat	ccc	528
Asn	Thr	Leu	Phe	Glu	Lys	Leu	Glu	Pro	Met	Ala	Ser	Ile	Leu	Asp	Pro	
				165				170						175		
ggc	gcg	agc	cag	agc	acg	acg	aac	ccg	atc	ttc	gga	atg	ccc	tcc	cct	576
Gly	Ala	Ser	Gln	Ser	Thr	Thr	Asn	Pro	Ile	Phe	Gly	Met	Pro	Ser	Pro	
			180				185						190			
ggc	agc	tca	aca	ccg	gtt	ggc	cag	ttg	ccg	ccg	gcg	gct	acc	cag	acc	624
Gly	Ser	Ser	Thr	Pro	Val	Gly	Gln	Leu	Pro	Pro	Ala	Ala	Thr	Gln	Thr	
			195			200							205			
ctc	ggc	caa	ctg	ggg	gag	atg	agc	ggc	ccg	atg	cag	cag	ctg	acc	cag	672
Leu	Gly	Gln	Leu	Gly	Glu	Met	Ser	Gly	Pro	Met	Gln	Gln	Leu	Thr	Gln	
	210				215						220					
ccg	ctg	cag	cag	gtg	acg	tcg	ttg	ttc	agc	cag	gtg	ggc	ggc	acc	ggc	720
Pro	Leu	Gln	Gln	Val	Thr	Ser	Leu	Phe	Ser	Gln	Val	Gly	Gly	Thr	Gly	
	225				230					235				240		
ggc	ggc	aac	cca	gcc	gac	gag	gaa	gcc	gcg	cag	atg	ggc	ctg	ctc	ggc	768
Gly	Gly	Asn	Pro	Ala	Asp	Glu	Glu	Ala	Ala	Gln	Met	Gly	Leu	Leu	Gly	
				245				250						255		
acc	agt	ccg	ctg	tcg	aac	cat	ccg	ctg	gct	ggg	tca	ggc	ccc	agc		816
Thr	Ser	Pro	Leu	Ser	Asn	His	Pro	Leu	Ala	Gly	Gly	Ser	Gly	Pro	Ser	
			260				265						270			
gcg	ggc	gcg	ggc	ctg	ctg	cgc	gcg	gag	tcg	cta	cct	ggc	gca	ggg	ggg	864
Ala	Gly	Ala	Gly	Leu	Leu	Arg	Ala	Glu	Ser	Leu	Pro	Gly	Ala	Gly	Gly	
			275			280							285			
tcg	ttg	acc	cgc	acg	ccg	ctg	atg	tct	cag	ctg	atc	gaa	aag	ccg	gtt	912
Ser	Leu	Thr	Arg	Thr	Pro	Leu	Met	Ser	Gln	Leu	Ile	Glu	Lys	Pro	Val	
	290				295					300						
gcc	ccc	tcg	gtg	atg	ccg	gcg	gct	gct	gcc	gga	tcg	tcg	gcg	acg	ggg	960
Ala	Pro	Ser	Val	Met	Pro	Ala	Ala	Ala	Ala	Gly	Ser	Ser	Ala	Thr	Gly	
	305				310					315				320		
ggc	gcc	gct	ccg	gtg	ggg	gcg	gga	gcg	atg	ggc	cag	ggg	gcg	caa	tcc	1008
Gly	Ala	Ala	Pro	Val	Gly	Ala	Gly	Ala	Met	Gly	Gln	Gly	Ala	Gln	Ser	
				325				330						335		
ggc	ggc	tcc	acc	agg	ccg	ggg	ctg	gtc	gcg	ccg	gca	ccg	ctc	gcg	cag	1056
Gly	Gly	Ser	Thr	Arg	Pro	Gly	Leu	Val	Ala	Pro	Ala	Pro	Leu	Ala	Gln	
			340				345						350			

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gag cgt gaa gaa gac gac gag gac gac tgg gac gaa gag gac gac tgg	1104
Glu Arg Glu Glu Asp Asp Glu Asp Asp Trp Asp Glu Glu Asp Asp Trp	
355 360 365	

tga	1107
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<210> SEQ ID NO 12  
 <211> LENGTH: 303  
 <212> TYPE: DNA  
 <213> ORGANISM: Mycobacterium tuberculosis  
 <220> FEATURE:  
 <221> NAME/KEY: CDS  
 <222> LOCATION: (1)...(300)

<400> SEQUENCE: 12

atg gca gag atg aag acc gat gcc gct acc ctc gcg cag gag gca ggt	48
Met Ala Glu Met Lys Thr Asp Ala Ala Thr Leu Ala Gln Glu Ala Gly	
1 5 10 15	

aat ttc gag cgg atc tcc gcc gac ctg aaa acc cag atc gac cag gtg	96
Asn Phe Glu Arg Ile Ser Gly Asp Leu Lys Thr Gln Ile Asp Gln Val	
20 25 30	

gag tcg acg gca ggt tcg ttg cag gcc cag tgg cgc gcc gcg gcg ggg	144
Glu Ser Thr Ala Gly Ser Leu Gln Gly Gln Trp Arg Gly Ala Ala Gly	
35 40 45	

acg gcc gcc cag gcc gcg gtg gtg cgc ttc caa gaa gca gcc aat aag	192
Thr Ala Ala Gln Ala Ala Val Val Arg Phe Gln Glu Ala Ala Asn Lys	
50 55 60	

cag aag cag gaa ctc gac gag atc tcg acg aat att cgt cag gcc gcc	240
Gln Lys Gln Glu Leu Asp Glu Ile Ser Thr Asn Ile Arg Gln Ala Gly	
65 70 75 80	

gtc caa tac tcg agg gcc gac gag gag cag cag cag gcg ctg tcc tcg	288
Val Gln Tyr Ser Arg Ala Asp Glu Glu Gln Gln Gln Ala Leu Ser Ser	
85 90 95	

caa atg gcc ttc tga	303
Gln Met Gly Phe	
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<210> SEQ ID NO 13  
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 <220> FEATURE:  
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 <222> LOCATION: (1)...(1998)

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Met Ala Ala Asp Tyr Asp Lys Leu Phe Arg Pro His Glu Gly Met Glu	
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gct ccg gac gat atg gca gcg cag ccg ttc ttc gac ccc agt gct tcg	96
Ala Pro Asp Asp Met Ala Ala Gln Pro Phe Phe Asp Pro Ser Ala Ser	
20 25 30	

ttt ccg ccg gcg ccc gca tcg gca aac cta ccg aag ccc aac gcc cag	144
Phe Pro Pro Ala Pro Ala Ser Ala Asn Leu Pro Lys Pro Asn Gly Gln	
35 40 45	

act ccg ccc ccg acg tcc gac gac ctg tcg gag ccg ttc gtg tcg gcc	192
Thr Pro Pro Pro Thr Ser Asp Asp Leu Ser Glu Arg Phe Val Ser Ala	
50 55 60	

ccg ccg ccg cca ccc cca ccc cca cct ccg cct ccg cca act ccg atg	240
Pro Pro Pro Pro Pro Pro Pro Pro Pro Pro Pro Pro Thr Pro Met	
65 70 75 80	

ccg atc gcc gca gga gag ccg ccc tcg ccg gaa ccg gcc gca tct aaa	288
Pro Ile Ala Ala Gly Glu Pro Pro Ser Pro Glu Pro Ala Ala Ser Lys	
85 90 95	

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cca ccc aca ccc ccc atg ccc atc gcc gga ccc gaa ccg gcc cca ccc Pro Pro Thr Pro Pro Met Pro Ile Ala Gly Pro Glu Pro Ala Pro Pro 100 105 110	336
aaa cca ccc aca ccc ccc atg ccc atc gcc gga ccc gaa ccg gcc cca Lys Pro Pro Thr Pro Pro Met Pro Ile Ala Gly Pro Glu Pro Ala Pro 115 120 125	384
ccc aaa cca ccc aca cct ccg atg ccc atc gcc gga cct gca ccc acc Pro Lys Pro Pro Thr Pro Pro Met Pro Ile Ala Gly Pro Ala Pro Thr 130 135 140	432
cca acc gaa tcc cag ttg gcg ccc ccc aga cca ccg aca cca caa acg Pro Thr Glu Ser Gln Leu Ala Pro Pro Arg Pro Pro Thr Pro Gln Thr 145 150 155 160	480
cca acc gga gcg ccg cag caa ccg gaa tca ccg gcg ccc cac gta ccc Pro Thr Gly Ala Pro Gln Gln Pro Glu Ser Pro Ala Pro His Val Pro 165 170 175	528
tcg cac ggg cca cat caa ccc ccg cgc acc gca cca gca ccg ccc tgg Ser His Gly Pro His Gln Pro Arg Arg Thr Ala Pro Ala Pro Pro Trp 180 185 190	576
gca aag atg cca atc ggc gaa ccc ccg ccc gct ccg tcc aga ccg tct Ala Lys Met Pro Ile Gly Glu Pro Pro Pro Ala Pro Ser Arg Pro Ser 195 200 205	624
gcg tcc ccg gcc gaa cca ccg acc ccg cct gcc ccc caa cac tcc cga Ala Ser Pro Ala Glu Pro Pro Thr Arg Pro Ala Pro Gln His Ser Arg 210 215 220	672
cgt gcg cgc ccg ggt cac cgc tat cgc aca gac acc gaa cga aac gtc Arg Ala Arg Arg Gly His Arg Tyr Arg Thr Asp Thr Glu Arg Asn Val 225 230 235 240	720
ggg aag gta gca act ggt cca tcc atc cag gcg ccg ctg ccg gca gag Gly Lys Val Ala Thr Gly Pro Ser Ile Gln Ala Arg Leu Arg Ala Glu 245 250 255	768
gaa gca tcc ggc gcg cag ctc gcc ccc gga acg gag ccc tcg cca gcg Glu Ala Ser Gly Ala Gln Leu Ala Pro Gly Thr Glu Pro Ser Pro Ala 260 265 270	816
ccg ttg ggc caa ccg aga tcg tat ctg gct ccg ccc acc cgc ccc gcg Pro Leu Gly Gln Pro Arg Ser Tyr Leu Ala Pro Pro Thr Arg Pro Ala 275 280 285	864
ccg aca gaa cct ccc ccc agc ccc tcg ccg cag cgc aac tcc ggt ccg Pro Thr Glu Pro Pro Pro Ser Pro Ser Pro Gln Arg Asn Ser Gly Arg 290 295 300	912
cgt gcc gag cga cgc gtc cac ccc gat tta gcc gcc caa cat gcc gcg Arg Ala Glu Arg Arg Val His Pro Asp Leu Ala Ala Gln His Ala Ala 305 310 315 320	960
gcg caa cct gat tca att acg gcc gca acc act ggc ggt cgt cgc cgc Ala Gln Pro Asp Ser Ile Thr Ala Ala Thr Thr Gly Gly Arg Arg Arg 325 330 335	1008
aag cgt gca gcg ccg gat ctc gac gcg aca cag aaa tcc tta agg ccg Lys Arg Ala Ala Pro Asp Leu Asp Ala Thr Gln Lys Ser Leu Arg Pro 340 345 350	1056
gcg gcc aag ggg ccg aag gtg aag aag gtg aag ccc cag aaa ccg aag Ala Ala Lys Gly Pro Lys Val Lys Lys Val Lys Pro Gln Lys Pro Lys 355 360 365	1104
gcc acg aag ccg ccc aaa gtg gtg tcg cag cgc ggc tgg cga cat tgg Ala Thr Lys Pro Pro Lys Val Val Ser Gln Arg Gly Trp Arg His Trp 370 375 380	1152
gtg cat gcg ttg acg cga atc aac ctg gcc ctg tca ccc gac gag aag Val His Ala Leu Thr Arg Ile Asn Leu Gly Leu Ser Pro Asp Glu Lys 385 390 395 400	1200
tac gag ctg gac ctg cac gct cga gtc cgc cgc aat ccc cgc ggg tcg Tyr Glu Leu Asp Leu His Ala Arg Val Arg Arg Asn Pro Arg Gly Ser	1248

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405	410	415	
tat cag atc gcc gtc gtc ggt ctc aaa ggt ggg gct ggc aaa acc acg Tyr Gln Ile Ala Val Val Gly Leu Lys Gly Gly Ala Gly Lys Thr Thr 420 425 430			1296
ctg aca gca gcg ttg ggg tcg acg ttg gct cag gtg cgg gcc gac cgg Leu Thr Ala Ala Leu Gly Ser Thr Leu Ala Gln Val Arg Ala Asp Arg 435 440 445			1344
atc ctg gct cta gac gcg gat cca ggc gcc gga aac ctc gcc gat cgg Ile Leu Ala Leu Asp Ala Asp Pro Gly Ala Gly Asn Leu Ala Asp Arg 450 455 460			1392
gta ggg cga caa tcg ggc gcg acc atc gct gat gtg ctt gca gaa aaa Val Gly Arg Gln Ser Gly Ala Thr Ile Ala Asp Val Leu Ala Glu Lys 465 470 475 480			1440
gag ctg tcg cac tac aac gac atc cgc gca cac act agc gtc aat gcg Glu Leu Ser His Tyr Asn Asp Ile Arg Ala His Thr Ser Val Asn Ala 485 490 495			1488
gtc aat ctg gaa gtg ctg ccg gca ccg gaa tac agc tcg gcg cag cgc Val Asn Leu Glu Val Leu Pro Ala Pro Glu Tyr Ser Ser Ala Gln Arg 500 505 510			1536
gcg ctc agc gac gcc gac tgg cat ttc atc gcc gat cct gcg tcg agg Ala Leu Ser Asp Ala Asp Trp His Phe Ile Ala Asp Pro Ala Ser Arg 515 520 525			1584
ttt tac aac ctc gtc ttg gct gat tgt ggg gcc ggc ttc ttc gac ccg Phe Tyr Asn Leu Val Leu Ala Asp Cys Gly Ala Gly Phe Phe Asp Pro 530 535 540			1632
ctg acc cgc ggc gtg ctg tcc acg gtg tcc ggt gtc gtg gtc gtg gca Leu Thr Arg Gly Val Leu Ser Thr Val Ser Gly Val Val Val Val Ala 545 550 555 560			1680
agt gtc tca atc gac ggc gca caa cag gcg tcg gtc gcg ttg gac tgg Ser Val Ser Ile Asp Gly Ala Gln Gln Ala Ser Val Ala Leu Asp Trp 565 570 575			1728
ttg cgc aac aac ggt tac caa gat ttg gcg agc cgc gca tgc gtg gtc Leu Arg Asn Asn Gly Tyr Gln Asp Leu Ala Ser Arg Ala Cys Val Val 580 585 590			1776
atc aat cac atc atg ccg gga gaa ccc aat gtc gca gtt aaa gac ctg Ile Asn His Ile Met Pro Gly Glu Pro Asn Val Ala Val Lys Asp Leu 595 600 605			1824
gtg cgg cat ttc gaa cag caa gtt caa ccc ggc cgg gtc gtg gtc atg Val Arg His Phe Glu Gln Gln Val Gln Pro Gly Arg Val Val Val Met 610 615 620			1872
ccg tgg gac agg cac att gcg gcc gga acc gag att tca ctc gac ttg Pro Trp Asp Arg His Ile Ala Ala Gly Thr Glu Ile Ser Leu Asp Leu 625 630 635 640			1920
ctc gac cct atc tac aag cgc aag gtc ctc gaa ttg gcc gca gcg cta Leu Asp Pro Ile Tyr Lys Arg Lys Val Leu Glu Leu Ala Ala Ala Leu 645 650 655			1968
tcc gac gat ttc gag agg gct gga cgt cgt tga Ser Asp Asp Phe Glu Arg Ala Gly Arg Arg 660 665			2001

&lt;210&gt; SEQ ID NO 14

&lt;211&gt; LENGTH: 1536

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&lt;221&gt; NAME/KEY: CDS

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&lt;400&gt; SEQUENCE: 14

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Ala Ala Arg Pro Ala Thr Thr Arg Val Thr Ile Leu Thr Gly Arg Arg	20	25	30	
atg acc gat ttg gta ctg cca gcg gcg gtg ccg atg gaa act tat att				144
Met Thr Asp Leu Val Leu Pro Ala Ala Val Pro Met Glu Thr Tyr Ile	35	40	45	
gac gac acc gtc gcg gtg ctt tcc gag gtg ttg gaa gac acg ccg gct				192
Asp Asp Thr Val Ala Val Leu Ser Glu Val Leu Glu Asp Thr Pro Ala	50	55	60	
gat gta ctc ggc ggc ttc gac ttt acc gcg caa ggc gtg tgg gcg ttc				240
Asp Val Leu Gly Gly Phe Asp Phe Thr Ala Gln Gly Val Trp Ala Phe	65	70	75	80
gct cgt ccc gga tgc ccg ccg ctg aag ctc gac cag tca ctc gat gac				288
Ala Arg Pro Gly Ser Pro Pro Leu Lys Leu Asp Gln Ser Leu Asp Asp	85	90	95	
gcc ggg gtg gtc gac ggg tca ctg ctg act ctg gtg tca gtc agt cgc				336
Ala Gly Val Val Asp Gly Ser Leu Leu Thr Leu Val Ser Val Ser Arg	100	105	110	
acc gag cgc tac cga ccg ttg gtc gag gat gtc atc gac gcg atc gcc				384
Thr Glu Arg Tyr Arg Pro Leu Val Glu Asp Val Ile Asp Ala Ile Ala	115	120	125	
gtg ctt gac gag tca cct gag ttc gac cgc acg gca ttg aat cgc ttt				432
Val Leu Asp Glu Ser Pro Glu Phe Asp Arg Thr Ala Leu Asn Arg Phe	130	135	140	
gtg ggg gcg gcg atc ccg ctt ttg acc gcg ccc gtc atc ggg atg gcg				480
Val Gly Ala Ala Ile Pro Leu Leu Thr Ala Pro Val Ile Gly Met Ala	145	150	155	160
atg cgg gcg tgg tgg gaa act ggg cgt agc ttg tgg tgg ccg ttg gcg				528
Met Arg Ala Trp Glu Thr Gly Arg Ser Leu Trp Trp Pro Leu Ala	165	170	175	
att ggc atc ctg ggg atc gct gtg ctg gta ggc agc ttc gtc gcg aac				576
Ile Gly Ile Leu Gly Ile Ala Val Leu Val Gly Ser Phe Val Ala Asn	180	185	190	
agg ttc tac cag agc ggc cac ctg gcc gag tgc cta ctg gtc acg acg				624
Arg Phe Tyr Gln Ser Gly His Leu Ala Glu Cys Leu Leu Val Thr Thr	195	200	205	
tat ctg ctg atc gca acc gcc gca gcg ctg gcc gtg ccg ttg ccg cgc				672
Tyr Leu Leu Ile Ala Thr Ala Ala Leu Ala Val Pro Leu Pro Arg	210	215	220	
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Gly Val Asn Ser Leu Gly Ala Pro Gln Val Ala Gly Ala Ala Thr Ala	225	230	235	240
gtg ctg ttt ttg acc ttg atg acg cgg ggc ggc cct cgg aag cgt cat				768
Val Leu Phe Leu Thr Leu Met Thr Arg Gly Gly Pro Arg Lys Arg His	245	250	255	
gag ttg gcg tcg ttt gcc gtg atc acc gct atc gcg gtc atc gcg gcc				816
Glu Leu Ala Ser Phe Ala Val Ile Thr Ala Ile Ala Val Ile Ala Ala	260	265	270	
gcc gct gcc ttc ggc tat gga tac cag gac tgg gtc ccc gcg ggg ggg				864
Ala Ala Ala Phe Gly Tyr Gly Tyr Gln Asp Trp Val Pro Ala Gly Gly	275	280	285	
atc gca ttc ggg ctg ttc att gtg acg aat gcg gcc aag ctg acc gtc				912
Ile Ala Phe Gly Leu Phe Ile Val Thr Asn Ala Ala Lys Leu Thr Val	290	295	300	
gcg gtc gcg cgg atc gcg ctg ccg ccg att ccg gta ccc ggc gaa acc				960
Ala Val Ala Arg Ile Ala Leu Pro Pro Ile Pro Val Pro Gly Glu Thr	305	310	315	320
gtg gac aac gag gag ttg ctc gat ccc gtc gcg acc ccg gag gct acc				1008



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Val Asp Asn Glu Glu Leu Leu Asp Pro Val Ala Thr Pro Glu Ala Thr	
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agc gaa gaa acc ccg acc tgg cag gcc atc atc gcg tgc gtg ccc gcg	1056
Ser Glu Glu Thr Pro Thr Trp Gln Ala Ile Ile Ala Ser Val Pro Ala	
340 345 350	
tcc gcg gtc cgg ctc acc gag cgc agc aaa ctg gcc aag caa ctt ctg	1104
Ser Ala Val Arg Leu Thr Glu Arg Ser Lys Leu Ala Lys Gln Leu Leu	
355 360 365	
atc gga tac gtc acg tgc ggc acc ctg att ctg gct gcc ggt gcc atc	1152
Ile Gly Tyr Val Thr Ser Gly Thr Leu Ile Leu Ala Ala Gly Ala Ile	
370 375 380	
gcg gtc gtg gtg cgc ggg cac ttc ttt gta cac agc ctg gtg gtc gcg	1200
Ala Val Val Val Arg Gly His Phe Phe Val His Ser Leu Val Val Ala	
385 390 395 400	
ggt ttg atc acg acc gtc tgc gga ttt cgc tgc cgg ctt tac gcc gag	1248
Gly Leu Ile Thr Thr Val Cys Gly Phe Arg Ser Arg Leu Tyr Ala Glu	
405 410 415	
cgc tgg tgt gcg tgg gcg ttg ctg gcg gcg acg gtc gcg att ccg acg	1296
Arg Trp Cys Ala Trp Ala Leu Leu Ala Ala Thr Val Ala Ile Pro Thr	
420 425 430	
ggt ctg acg gcc aaa ctc atc atc tgg tac ccg cac tat gcc tgg ctg	1344
Gly Leu Thr Ala Lys Leu Ile Ile Trp Tyr Pro His Tyr Ala Trp Leu	
435 440 445	
ttg ttg agc gtc tac ctc acg gta gcc ctg gtt gcg ctc gtg gtg gtc	1392
Leu Leu Ser Val Tyr Leu Thr Val Ala Leu Val Ala Leu Val Val Val	
450 455 460	
ggg tgc atg gct cac gtc cgg cgc gtt tca ccg gtc gta aaa cga act	1440
Gly Ser Met Ala His Val Arg Arg Val Ser Pro Val Val Lys Arg Thr	
465 470 475 480	
ctg gaa ttg atc gac ggc gcc atg atc gct gcc atc att ccc atg ctg	1488
Leu Glu Leu Ile Asp Gly Ala Met Ile Ala Ala Ile Ile Pro Met Leu	
485 490 495	
ctg tgg atc acc ggg gtg tac gac acg gtc cgc aat atc cgg ttc	1533
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tga	1536
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gcg aaa ttg gcc ggc ctc gtt ttt ccg cag cct ccg gcg ccg atc gcg	96
Ala Lys Leu Ala Gly Leu Val Phe Pro Gln Pro Pro Ala Pro Ile Ala	
20 25 30	
gtc agc gga acg gat tgc gtg gta gca gca atc aac gag acc atg cca	144
Val Ser Gly Thr Asp Ser Val Val Ala Ala Ile Asn Glu Thr Met Pro	
35 40 45	
agc atc gaa tgc ctg gtc agt gac ggg ctg ccc ggc gtg aaa gcc gcc	192
Ser Ile Glu Ser Leu Val Ser Asp Gly Leu Pro Gly Val Lys Ala Ala	
50 55 60	
ctg act cga aca gca tcc aac atg aac gcg gcg gcg gac gtc tat gcg	240
Leu Thr Arg Thr Ala Ser Asn Met Asn Ala Ala Ala Asp Val Tyr Ala	
65 70 75 80	

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aag acc gat cag tca ctg gga acc agt ttg agc cag tat gca ttc ggc Lys Thr Asp Gln Ser Leu Gly Thr Ser Leu Ser Gln Tyr Ala Phe Gly	288
85 90 95	
tcg tcg ggc gaa ggc ctg gct ggc gtc gcc tcg gtc ggt ggt cag cca Ser Ser Gly Glu Gly Leu Ala Gly Val Ala Ser Val Gly Gly Gln Pro	336
100 105 110	
agt cag gct acc cag ctg ctg agc aca ccc gtg tca cag gtc acg acc Ser Gln Ala Thr Gln Leu Leu Ser Thr Pro Val Ser Gln Val Thr Thr	384
115 120 125	
cag ctc ggc gag acg gcc gct gag ctg gca ccc cgt gtt gtt gcg acg Gln Leu Gly Glu Thr Ala Ala Glu Leu Ala Pro Arg Val Val Ala Thr	432
130 135 140	
gtg ccg caa ctc gtt cag ctg gct ccg cac gcc gtt cag atg tcg caa Val Pro Gln Leu Val Gln Leu Ala Pro His Ala Val Gln Met Ser Gln	480
145 150 155 160	
aac gca tcc ccc atc gct cag acg atc agt caa acc gcc caa cag gcc Asn Ala Ser Pro Ile Ala Gln Thr Ile Ser Gln Thr Ala Gln Gln Ala	528
165 170 175	
gcc cag agc gcg cag ggc ggc agc ggc cca atg ccc gca cag ctt gcc Ala Gln Ser Ala Gln Gly Gly Ser Gly Pro Met Pro Ala Gln Leu Ala	576
180 185 190	
agc gct gaa aaa ccg gcc acc gag caa gcg gag ccg gtc cac gaa gtg Ser Ala Glu Lys Pro Ala Thr Glu Gln Ala Glu Pro Val His Glu Val	624
195 200 205	
aca aac gac gat cag ggc gac cag ggc gac gtg cag ccg gcc gag gtc Thr Asn Asp Asp Gln Gly Asp Gln Gly Asp Val Gln Pro Ala Glu Val	672
210 215 220	
gtt gcc gcg gca cgt gac gaa ggc gcc ggc gca tca ccg ggc cag cag Val Ala Ala Ala Arg Asp Glu Gly Ala Gly Ala Ser Pro Gly Gln Gln	720
225 230 235 240	
ccc ggc ggg ggc gtt ccc gcg caa gcc atg gat acc gga gcc ggt gcc Pro Gly Gly Gly Val Pro Ala Gln Ala Met Asp Thr Gly Ala Gly Ala	768
245 250 255	
cgc cca gcg gcg agt ccg ctg gcg gcc ccc gtc gat ccg tcg act ccg Arg Pro Ala Ala Ser Pro Leu Ala Ala Pro Val Asp Pro Ser Thr Pro	816
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gca ccc tca aca acc aca acg ttg tag Ala Pro Ser Thr Thr Thr Thr Leu	843
275 280	
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ccg ggc ggc tgg gtg gaa gcc gat gaa gac act ttc tat gac ccg gcc Pro Gly Gly Trp Val Glu Ala Asp Glu Asp Thr Phe Tyr Asp Arg Ala	96
20 25 30	
cag gaa tat agc cag gtt ttg caa agg gtc acc gat gta ttg gac acc Gln Glu Tyr Ser Gln Val Leu Arg Val Thr Asp Val Leu Asp Thr	144
35 40 45	
tgc cgc cag cag aaa ggc cac gtc ttc gaa ggc ggc cta tgg tcc ggc Cys Arg Gln Gln Lys Gly His Val Phe Glu Gly Gly Leu Trp Ser Gly	192
50 55 60	

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cat att gcc ggg ttg att gag caa gct aaa tcc gat atc ggc aat aat His Ile Ala Gly Leu Ile Glu Gln Ala Lys Ser Asp Ile Gly Asn Asn 100 105 110	336
gtg gat ggc gct caa cgg gag atc gat atc ctg gag aat gac cct agc Val Asp Gly Ala Gln Arg Glu Ile Asp Ile Leu Glu Asn Asp Pro Ser 115 120 125	384
ctg gat gct gat gag cgc cat acc gcc atc aat tca ttg gtc acg gcg Leu Asp Ala Asp Glu Arg His Thr Ala Ile Asn Ser Leu Val Thr Ala 130 135 140	432
acg cat ggg gcc aat gtc agt ctg gtc gcc gag acc gct gag cgg gtg Thr His Gly Ala Asn Val Ser Leu Val Ala Glu Thr Ala Glu Arg Val 145 150 155 160	480
ctg gaa tcc aag aat tgg aaa cct ccg aag aac gca ctc gag gat ttg Leu Glu Ser Lys Asn Trp Lys Pro Pro Lys Asn Ala Leu Glu Asp Leu 165 170 175	528
ctt cag cag aag tgc ccg cca ccc cca gac gtg cct acc ctg gtc gtg Leu Gln Gln Lys Ser Pro Pro Pro Asp Val Pro Thr Leu Val Val 180 185 190	576
cca tcc ccg ggc aca ccg ggc aca ccg gga acc ccg atc acc ccg gga Pro Ser Pro Gly Thr Pro Gly Thr Pro Gly Thr Pro Ile Thr Pro Gly 195 200 205	624
acc ccg atc acc ccg gga acc cca atc aca ccc atc ccg gga gcg ccg Thr Pro Ile Thr Pro Gly Thr Pro Ile Thr Pro Ile Pro Gly Ala Pro 210 215 220	672
gta act ccg atc aca cca acg ccc ggc act ccc gtc acg ccg gtg acc Val Thr Pro Ile Thr Pro Thr Pro Gly Thr Pro Val Thr Pro Val Thr 225 230 235 240	720
ccg ggc aag ccg gtc acc ccg gtg acc ccg gtc aaa ccg ggc aca cca Pro Gly Lys Pro Val Thr Pro Val Thr Pro Val Lys Pro Gly Thr Pro 245 250 255	768
ggc gag cca acc ccg atc acg ccg gtc acc ccc ccg gtc gcc ccg gcc Gly Glu Pro Thr Pro Ile Thr Pro Val Thr Pro Pro Val Ala Pro Ala 260 265 270	816
aca ccg gca acc ccg gcc acg ccc gtt acc cca gct ccc gct cca cac Thr Pro Ala Thr Pro Ala Thr Pro Val Thr Pro Ala Pro Ala Pro His 275 280 285	864
ccg cag ccg gct ccg gca ccg ggc cca tgc cct ggg ccc cag ccg gtt Pro Gln Pro Ala Pro Ala Pro Ala Pro Ser Pro Gly Pro Gln Pro Val 290 295 300	912
aca ccg gcc act ccc ggt ccg tct ggt cca gca aca ccg ggc acc cca Thr Pro Ala Thr Pro Gly Pro Ser Gly Pro Ala Thr Pro Gly Thr Pro 305 310 315 320	960
ggg ggc gag ccg gcg ccg cac gtc aaa ccc gcg gcg ttg gcg gag caa Gly Gly Glu Pro Ala Pro His Val Lys Pro Ala Ala Leu Ala Glu Gln 325 330 335	1008
cct ggt gtg ccg ggc cag cat gcg ggc ggg ggg acg cag tgc ggg cct Pro Gly Val Pro Gly Gln His Ala Gly Gly Gly Thr Gln Ser Gly Pro 340 345 350	1056
gcc cat gcg gac gaa tcc gcc gcg tgc gtg acg ccg gct gcg gcg tcc Ala His Ala Asp Glu Ser Ala Ala Ser Val Thr Pro Ala Ala Ala Ser 355 360 365	1104
ggt gtc ccg ggc gca ccg gcg gcg gcc gcc gcg ccg agc ggt acc gcc Gly Val Pro Gly Ala Arg Ala Ala Ala Ala Pro Ser Gly Thr Ala 370 375 380 385 390 395	1152

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370					375					380							
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gcg	ggg	tcg	cat	gct	gcc	act	ggg	cgg	gcg	ccg	gtg	gct	acc	tcg	gac	1248	
Ala	Gly	Ser	His	Ala	Ala	Thr	Gly	Arg	Ala	Pro	Val	Ala	Thr	Ser	Asp		
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aag	gcg	gcg	gca	ccg	agc	acg	cgg	gcg	gcc	tcg	gcg	cgg	acg	gca	cct	1296	
Lys	Ala	Ala	Ala	Pro	Ser	Thr	Arg	Ala	Ala	Ser	Ala	Arg	Thr	Ala	Pro		
					420					425					430		
cct	gcc	cgc	ccg	ccg	tcg	acc	gat	cac	atc	gac	aaa	ccc	gat	cgc	agc	1344	
Pro	Ala	Arg	Pro	Pro	Ser	Thr	Asp	His	Ile	Asp	Lys	Pro	Asp	Arg	Ser		
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gag	tct	gca	gat	gac	ggt	acg	ccg	gtg	tcg	atg	atc	ccg	gtg	tcg	gcg	1392	
Glu	Ser	Ala	Asp	Asp	Gly	Thr	Pro	Val	Ser	Met	Ile	Pro	Val	Ser	Ala		
					450					455					460		
gct	cgg	gcg	gca	cgc	gac	gcc	gcc	act	gca	gct	gcc	agc	gcc	cgc	cag	1440	
Ala	Arg	Ala	Ala	Arg	Asp	Ala	Ala	Thr	Ala	Ala	Ala	Ser	Ala	Arg	Gln		
					465					470					475		480
cgt	ggc	cgc	ggt	gat	gcg	ctg	cgg	ttg	gcg	cga	cgc	atc	gcg	gcg	gcg	1488	
Arg	Gly	Arg	Gly	Asp	Ala	Leu	Arg	Leu	Ala	Arg	Arg	Ile	Ala	Ala	Ala		
					485					490					495		
ctc	aac	gcg	tcc	gac	aac	aac	gcg	ggc	gac	tac	ggg	ttc	ttc	tgg	atc	1536	
Leu	Asn	Ala	Ser	Asp	Asn	Asn	Ala	Gly	Asp	Tyr	Gly	Phe	Phe	Trp	Ile		
					500					505					510		
acc	gcg	gtg	acc	acc	gac	ggt	tcc	atc	gtc	gtg	gcc	aac	agc	tat	ggg	1584	
Thr	Ala	Val	Thr	Thr	Asp	Gly	Ser	Ile	Val	Val	Ala	Asn	Ser	Tyr	Gly		
					515					520					525		
ctg	gcc	tac	ata	ccc	gac	ggg	atg	gaa	ttg	ccg	aat	aag	gtg	tac	ttg	1632	
Leu	Ala	Tyr	Ile	Pro	Asp	Gly	Met	Glu	Leu	Pro	Asn	Lys	Val	Tyr	Leu		
					530					535					540		
gcc	agc	gcg	gat	cac	gca	atc	ccg	ggt	gac	gaa	att	gca	cgc	tgt	gcc	1680	
Ala	Ser	Ala	Asp	His	Ala	Ile	Pro	Val	Asp	Glu	Ile	Ala	Arg	Cys	Ala		
					545					550					555		560
acc	tac	ccg	ggt	ttg	gcc	gtg	caa	gcc	tgg	gcg	gct	ttc	cac	gac	atg	1728	
Thr	Tyr	Pro	Val	Leu	Ala	Val	Gln	Ala	Trp	Ala	Ala	Phe	His	Asp	Met		
					565					570					575		
acg	ctg	cgg	gcg	gtg	atc	ggt	acc	gcg	gag	cag	ttg	gcc	agt	tcg	gat	1776	
Thr	Leu	Arg	Ala	Val	Ile	Gly	Thr	Ala	Glu	Gln	Leu	Ala	Ser	Ser	Asp		
					580					585					590		
ccc	ggt	gtg	gcc	aag	att	gtg	ctg	gag	cca	gat	gac	att	ccg	gag	agc	1824	
Pro	Gly	Val	Ala	Lys	Ile	Val	Leu	Glu	Pro	Asp	Asp	Ile	Pro	Glu	Ser		
					595					600					605		
ggc	aaa	atg	acg	ggc	cgg	tcg	cgg	ctg	gag	gtc	gtc	gac	ccc	tcg	gcg	1872	
Gly	Lys	Met	Thr	Gly	Arg	Ser	Arg	Leu	Glu	Val	Val	Asp	Pro	Ser	Ala		
					610					615					620		
gcg	gct	cag	ctg	gcc	gac	act	acc	gat	cag	cgt	ttg	ctc	gac	ttg	ttg	1920	
Ala	Ala	Gln	Leu	Ala	Asp	Thr	Thr	Asp	Gln	Arg	Leu	Leu	Asp	Leu	Leu		
					625					630					635		640
ccg	ccg	gcg	ccg	gtg	gat	gtc	aat	cca	ccg	ggc	gat	gag	cgg	cac	atg	1968	
Pro	Pro	Ala	Pro	Val	Asp	Val	Asn	Pro	Pro	Gly	Asp	Glu	Arg	His	Met		
					645					650					655		
ctg	tgg	ttc	gag	ctg	atg	aag	ccc	atg	acc	agc	acc	gct	acc	ggc	cgc	2016	
Leu	Trp	Phe	Glu	Leu	Met	Lys	Pro	Met	Thr	Ser	Thr	Ala	Thr	Gly	Arg		
					660					665					670		
gag	gcc	gct	cat	ctg	cgg	gcg	ttc	cgg	gcc	tac	gct	gcc	cac	tca	cag	2064	
Glu	Ala	Ala	His	Leu	Arg	Ala	Phe	Arg	Ala	Tyr	Ala	Ala	His	Ser	Gln		
					675					680					685		
gag	att	gcc	ctg	cac	caa	gcg	cac	act	gcg	act	gac	gcg	gcc	gtc	cag	2112	

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Glu	Ile	Ala	Leu	His	Gln	Ala	His	Thr	Ala	Thr	Asp	Ala	Ala	Val	Gln	
690						695					700					
cgt	gtg	gcc	gtc	gcg	gac	tgg	ctg	tac	tgg	caa	tac	gtc	acc	ggg	ttg	2160
Arg	Val	Ala	Val	Ala	Asp	Trp	Leu	Tyr	Trp	Gln	Tyr	Val	Thr	Gly	Leu	
705					710					715					720	
ctc	gac	cgg	gcc	ctg	gcc	gcc	gca	tgc	tga							2190
Leu	Asp	Arg	Ala	Leu	Ala	Ala	Ala	Cys								
					725											

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What is claimed:

1. A method of in vitro diagnosis that discriminates between infection by *Mycobacterium tuberculosis*-complex and vaccination by Bacille Calmette Guerin (BCG) strain of *Mycobacterium bovis* comprising:

providing a population of cells comprising CD4 T lymphocytes from a subject;

contacting cells of the population with at least two different antigens, wherein the antigens are isolated polypeptides of the *Mycobacterium tuberculosis*-complex that are not encoded by BCG, including at least one isolated polypeptide selected from the group consisting of (i) a first amino acid sequence comprising the sequence of MTBN4 (SEQ ID NO: 4), (ii) a second amino acid sequence that is an antigenic segment of MTBN4 and (iii) a third amino acid sequence that is identical to said first or second amino acid sequence but that has conservative substitutions and that retains *Mycobacterium tuberculosis*-complex specific antigenic properties; and

determining whether or not there has been an immune response to said at least two different antigens, wherein CD4 T lymphocytes from a subject that has been infected by *Mycobacterium tuberculosis*-complex

respond to said at least two different antigens, and CD4 T lymphocytes from a subject vaccinated with the BCG strain of *Mycobacterium bovis* but not infected by *Mycobacterium tuberculosis*-complex do not respond to said at least two different antigens.

2. The method of claim 1, wherein said at least one isolated polypeptide comprises said second or third amino acid sequence.

3. The method of claim 1, wherein the step of contacting is contacting said cells with a composition containing said at least two different antigens.

4. The method of claim 3, wherein the determining step comprises testing for production of at least one cytokine.

5. The method of claim 4, wherein the at least one cytokine includes interferon- $\gamma$  (IFN- $\gamma$ ).

6. The method of claim 1, wherein the determining step comprises testing for production of at least one cytokine.

7. The method of claim 6, wherein the at least one cytokine includes interferon- $\gamma$  (IFN- $\gamma$ ).

8. The method of claim 1, wherein the isolated polypeptides of the *Mycobacterium tuberculosis*-complex are encoded within the RD1, RD2, and RD3 regions.

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